(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization International Bureau



(43) International Publication Date 3 May 2001 (03.05.2001)

PCT

(10) International Publication Number WO 01/30991 A3

- (51) International Patent Classification7: C07K 14/705. C12Q 1/68
- (21) International Application Number: PCT/US00/23021
- (22) International Filing Date: 22 August 2000 (22.08.2000)
- (25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

09/426,290

25 October 1999 (25.10.1999)

(63) Related by continuation (CON) or continuation-in-part (CIP) to earlier application:

US

09/426,290 (CIP)

Filed on

25 October 1999 (25.10.1999)

- (71) Applicant (for all designated States except US): DECODE GENETICS EHF. [IS/IS]: Lynghals 1, IS-110 Reykjavík (IS).
- (72) Inventors; and
- (75) Inventors/Applicants (for US only): OLAFSDOTTIR, Berglind, Ran [IS/IS]; Eskihlid 15, IS-105 Reykjavik (IS). GULCHER, Jeffrey [US/US]; Unit M, 130 South Canal Street, Chicago, IL 60606 (US).

- (74) Agents: CARROLL, Alice, O. et al.; Hamilton, Brook. Smith & Reynolds, P.C., 530 Virginia Road, P.O. Box 9133, Concord, MA 01742-9133 (US).
- (81) Designated States (national): AE. AG. AL. AM. AT. AU. AZ. BA, BB, BG, BR. BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO. NZ, PL. PT. RO, RU, SD. SE, SG, SI, SK, SL, TJ, TM. TR. TT, TZ, UA. UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW. MZ. SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT. LU. MC, NL, PT. SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

with international search report

(88) Date of publication of the international search report: 6 December 2001

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: HUMAN NARCOLEPSY GENE

(57) Abstract: The gene for hypocretin (orexin) receptor 2 (HCRTR2), which is associated with narcolepsy, is disclosed. Also described are methods of diagnosis of narcolepsy, pharmaceutical compositions comprising nucleic acids comprising the HCRTR2 gene, as well as methods of therapy of narcolepsy.

Internat. J Application No PCT/US 00/23021

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C07K14/705 C12Q1/68

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols) IPC 7 $\,$ C07K $\,$ C12Q

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, WPI Data, BIOSIS

C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to daim No.		
Υ	WO 96 34877 A (HUMAN GENOME SCIENCES INC.; LI YI (US); ROSEN CRAIG A (US); SOPPET) 7 November 1996 (1996-11-07) the whole document	1-7		
Y	LIN LING ET AL: "The sleep disorder canine narcolepsy is caused by a mutation in the hypocretin (orexin) receptor 2 gene" CELL,CELL PRESS, CAMBRIDGE, NA,US, vol. 98, no. 3, 6 August 1999 (1999-08-06), pages 365-376, XP002153571 ISSN: 0092-8674 abstract; figure 6	1-7		

Further documents are listed in the continuation of box C.	Patent family members are listed in annex.
Special categories of cited documents: A' document defining the general state of the art which is not considered to be of particular relevance E' earlier document but published on or after the international filing date L' document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) O' document reterring to an oral disclosure, use, exhibition or other means P' document published prior to the international filing date but later than the priority date claimed	'T' later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention 'X' document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone 'Y' document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. '&' document member of the same patent family
Date of the actual completion of the international search 22 March 2001	Date of mailing of the international search report
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Authorized officer Hardon, E

1

Internat. J Application No PCT/US 00/23021

	TO BE THE CONTRIBUTION OF	PCT/US 00/23021
C.(Continua Category °	ontion) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
	TP PP P	rioles dill to Ciditi (40.
Y	SAKURAI T ET AL: "Oxerins and oxerin receptors: A family of hypothalamic neuropeptides and G Protein-coupled receptors that regulate feeding behaviour" CELL, CELL PRESS, CAMBRIDGE, NA, US, vol. 92, 20 February 1998 (1998-02-20), pages 573-585, XP002105412 ISSN: 0092-8674 page 585, column 2; figure 2	1-7
Y	ALDRICH, MICHAEL S. ET AL: "Narcolepsy and the hypocretin receptor 2 gene" NEURON (1999), 23(4), 625-626 , 1999, XP000973742 the whole document	1-7
Y	SIEGEL, JEROME M.: "Narcolepsy: A key role for hypocretins (orexins)" CELL (CAMBRIDGE, MASS.) (1999), 98(4), 409-412, 20 August 1999 (1999-08-20), XP000941943 the whole document	1-7
A	LECEA L ET AL: "The hypocretins: hypothalamus-specific peptides with neuroexcitatory activity" PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA,NATIONAL ACADEMY OF SCIENCE. WASHINGTON,US, vol. 95, January 1998 (1998-01), pages 322-327, XP002105411 ISSN: 0027-8424 the whole document	1-7
T	PEYRON CHRISTELLE ET AL: "A mutation in a case of early onset narcolepsy and a generalized absence of hypocretin peptides in human narcoleptic brains" NATURE MEDICINE, NATURE PUBLISHING, CO,US, vol. 6, no. 9, September 2000 (2000-09), pages 991-997, XP002153570 ISSN: 1078-8956	1-7
	·	

Inten ational application No. PCT/US 00/23021

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)
This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1. X Claims Nos.: 7 because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy
2. Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)
This International Searching Authority found multiple inventions in this international application, as follows:
As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remark on Protest The additional search fees were accompanied by the applicant's protest. No protest accompanied the payment of additional search fees.

· · · · · · · · · · · · · · · · · · ·	nformation on patent family m		PCT/US	00/23021
Patent document dted in search report	Publication date	Patent fam member(s		Publication date
WO 9634877 A	07-11-1996	AU 715 AU 2470 EP 0828	036 A 286 B 795 A 751 A	07-11-1996 20-01-2000 21-11-1996 18-03-1998 18-05-1999
		·		

(19) World Intellectual Property Organization International Bureau



| 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 1886 | 188

(43) International Publication Date 3 May 2001 (03.05.2001)

PCT

(10) International Publication Number WO 01/30991 A2

(51) International Patent Classification7: C

C12N 15/00

(21) International Application Number: PCT/US00/23021

(22) International Filing Date: 22 August 2000 (22.08.2000)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

09/426,290

25 October 1999 (25.10.1999) US

(63) Related by continuation (CON) or continuation-in-part (CIP) to earlier application:

US Filed on 09/426,290 (CIP) 25 October 1999 (25.10.1999)

(71) Applicant (for all designated States except US): DECODE GENETICS EHF. [IS/IS]; Lynghals 1, IS-110 Reykjavik

(21)

(72) Inventors; and

(75) Inventors/Applicants (for US only): OLAFSDOTTIR, Berglind, Ran [IS/IS]; Eskihlid 15, IS-105 Reykjavik (IS). GULCHER, Jeffrey [US/US]; Unit M, 130 South Canal Street, Chicago, IL 60606 (US). 74) Agents: CARROLL, Alice, O. et al.; Hamilton, Brook, Smith & Reynolds, P.C., Two Militia Drive, Lexington, MA 02421 (US).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

 Without international search report and to be republished upon receipt of that report.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: HUMAN NARCOLEPSY GENE

(57) Abstract: The gene for hypocretin (orexin) receptor 2 (HCRTR2), which is associated with narcolepsy, is disclosed. Also described are methods of diagnosis of narcolepsy, pharmaceutical compositions comprising nucleic acids comprising the HCRTR2 gene, as well as methods of therapy of narcolepsy.

-1-

HUMAN NARCOLEPSY GENE

RELATED APPLICATION

5

This application is a Continuation-in-Part of U.S. Serial No. 09/426,290, filed October 25, 1999, the entire teachings of which are incorporated herein by reference.

BACKGROUND OF THE INVENTION

Narcolepsy, a disorder which affects approximately 1 in 2,000 individuals, is characterized by daytime sleepiness, sleep fragmentation, and symptoms of abnormal rapid eye movement (REM) sleep that include cataplexy (loss of muscle 10 tone), sleep paralysis, and hypnagogic hallucinations (Aldrich, M.S., Neurology 42:34-43 (1992); Siegel, J.M., Cell 98:409-412 (1999)). In humans, susceptibility to narcolepsy has been associated with a specific human leukocyte antigen (HLA) alleles, including DQB1*0602 (Mignot, E., Neurology 50:S16-22 (1998); Kadotani, H. et al., Genome Res. 8:427-434 (1998); Faraco, J. et al., J. Hered. 90:129-132 15 (1999)); however, attempts to verify narcolepsy as an autoimmune disorder have failed (Mignot, E. et al., Adv. Neuroimmunol. 5:23-37 (1995); Mignot, E., Curr. Opin. Pulm. Med. 2:482-487 (1996)). In a canine model of narcolepsy, the disorder is transmitted as an autosomal recessive trait, canarc-1 (Foutz, A.S. et al., Sleep 1:413-421 91979); Baker, T.L. and Dement, W.C., Brain Mechanisms of Sleep (D.J. McGinty et al., eds., New York: Raven Press, pp. 199-233 (1985)). The possibility of linkage between canarc-1 and the canine major histocompatibility complex has been excluded (Mignot, E. et al., Proc. Natl. Acad. Sci. USA 88:3475-3478 (1991)).

15

25

A mutation in the hypocretin (orexin) receptor 2 gene in canines has been identified in narcolepsy (Lin, L. et al., Cell 98:365-376 (1999)); Hypocrexins/orexins (orexin-A and -B) are neuropeptides associated with regulation of food consumption (de Lecea, L., et al., Proc. natl. Acad. Sci. USA 95:322-327 (1998); Sakurai, T. et al., Cell 92:573-585 (1998)) as well as other possible functions (Peyron, C. et al., J. Neurosci. 18:9996-10015 (1998)). Human cDNA of receptors for orexins have been cloned (Sakurai, T. et al., Cell 92:573-585 (1998)), however, full human genes for the orexin receptors have not yet been identified.

Diagnosis of narcolepsy is difficult, as it is necessary to distinguish narcolepsy from other conditions such as chronic fatigue syndrome or other sleep 10 disorders (Ambrogetti, A. and Olson, L.C., Med. J. Aust. 160:426-429 (1994); Aldrich, M.S., Neurology 50:S2-7 (1998)). Methods of diagnosing narcolepsy based on specific criteria would facilitate identification of the disease, reduce the time and expense associated with diagnosis, and expedite commencement of treatment.

SUMMARY OF THE INVENTION

As described herein, a full gene for the human hypocretin (orexin) receptor 2 (HCRTR2) has been identified. The sequence of the HCRTR2 gene as described herein is shown in Figure 1 (SEQ ID NO: 1). Accordingly, this invention pertains to an isolated nucleic acid molecule containing the HCRTR2 gene. The invention also relates to DNA constructs comprising the nucleic acid molecules described herein operatively linked to a regulatory sequence, and to recombinant host cells, such as bacterial cells, fungal cells, plant cells, insect cells and mammalian cells, comprising the nucleic acid molecules described herein operatively linked to a regulatory sequence. The invention also pertains to methods of diagnosing narcolepsy in an individual. The methods include detecting the presence of a mutation in the HCRTR2 gene. The invention additionally pertains to pharmaceutical compositions comprising the HCRTR2 nucleic acids of the invention. The invention further pertains to methods of treating narcolepsy, by administering HCRTR2 nucleic acids

of the invention or compositions comprising the HCRTR2 nucleic acids. The methods of the invention allow the accurate diagnosis of narcolepsy and reduce the need for time-consuming and expensive sleep laboratory assessments.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1A to Fig. 1AY depict the sequence of the human orexin receptor 2 gene (SEQ ID NO:1) and the encoded receptor (SEQ ID NO:2).

The foregoing and other objects, features and advantages of the invention will be apparent from the following more particular description of preferred embodiments of the invention, as illustrated in the accompanying drawings

10 DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to a human hypocretin (orexin) receptor 2 (HCRTR2) gene, and the relationship of the gene to narcolepsy. As described herein, Applicants have isolated the HCRTR2 gene. The gene and its products are implicated in the pathogenesis of narcolepsy, as mutations in a closely related receptor, hypocretin (orexin) receptor 2, have been associated with the presence of narcolepsy in a well-established canine model of narcolepsy (Lin, L. et al., Cell 98:365-376 (1999)).

NUCLEIC ACIDS OF THE INVENTION

Accordingly, the invention pertains to an isolated nucleic acid molecule

containing the human HCRTR2 gene. The term, "HCRTR2 gene," refers to an
isolated genomic nucleic acid molecule that encodes the human hypocretin (orexin)
receptor 2. As used herein, the term, "genomic nucleic acid molecule" indicates that
the nucleic acid molecule contains introns and exons as are found in genomic DNA
(i.e., not cDNA). The nucleic acid molecules can be double-stranded or singlestranded; single stranded nucleic acid molecules can be either the coding (sense)
strand or the non-coding (antisense) strand. The nucleic acid molecule can
additionally contain a marker sequence, for example, a nucleotide sequence which
encodes a polypeptide, to assist in isolation or purification of the polypeptide. Such

5

10

20

25

30

-4-

sequences include, but are not limited to, those which encode a glutathione-S-transferase (GST) fusion protein and those which encode a hemagglutinin A (HA) peptide marker from influenza. In a preferred embodiment, the nucleic acid molecule has the sequence shown in the Figure (SEQ ID NO:1).

As used herein, an "isolated" or "substantially pure" gene or nucleic acid molecule is intended to mean a gene which is not flanked by nucleotide sequences which normally (in nature) flank the gene (as in other genomic sequences). Thus, an isolated gene can include a gene which is synthesized chemically or by recombinant means. Thus, recombinant DNA contained in a vector are included in the definition of "isolated" as used herein. Also, isolated nucleotide sequences include recombinant DNA molecules in heterologous host cells, as well as partially or substantially purified DNA molecules in solution. Such isolated nucleotide sequences are useful in the manufacture of the encoded protein, as probes for isolating homologous sequences (e.g., from other mammalian species), for gene mapping (e.g., by *in situ* hybridization with chromosomes), or for detecting expression of the HCRTR2 gene in tissue (e.g., human tissue), such as by Northern blot analysis.

The present invention also encompasses variations of the nucleic acid sequences of the invention. Such variations can be naturally-occurring, such as in the case of allelic variation, or non-naturally-occurring, such as those induced by various mutagens and mutagenic processes. Intended variations include, but are not limited to, addition, deletion and substitution of one or more nucleotides which can result in conservative or non-conservative amino acid changes, including additions and deletions. Preferably, the nucleotide or amino acid variations are silent or conserved; that is, they do not alter the characteristics or activity of the hypocretin (orexin) receptor 2.

Other alterations of the nucleic acid molecules of the invention can include, for example, labeling, methylation, internucleotide modifications such as uncharged linkages (e.g., methyl phosphonates, phosphotriesters, phosphoamidates, carbamates), charged linkages (e.g., phosphorothioates, phosphorodithioates), pendent moieties (e.g., polypeptides), intercalators (e.g., acridine, psoralen),

10

15

20

25

30

chelators, alkylators, and modified linkages (e.g., alpha anomeric nucleic acids).

Also included are synthetic molecules that mimic nucleic acid molecules in the ability to bind to a designated sequences via hydrogen bonding and other chemical interactions. Such molecules include, for example, those in which peptide linkages substitute for phosphate linkages in the backbone of the molecule.

-5-

PCT/US00/23021

The invention also relates to fragments of the isolated nucleic acid molecules described herein. The term "fragment" is intended to encompass a portion of a nucleic acid sequence described herein which is from at least about 25 contiguous nucleotides to at least about 50 contiguous nucleotides or longer in length. One or more introns can also be present. Such fragments are useful as probes, e.g., for diagnostic methods, as described below and also as primers or probes. Particularly preferred primers and probes selectively hybridize to a nucleic acid molecule containing the HCRTR2 gene described herein.

The invention also pertains to nucleic acid molecules which hybridize under high stringency hybridization conditions, such as for selective hybridization, to a nucleotide sequence described herein (e.g., nucleic acid molecules which specifically hybridize to a nucleic acid containing the HCRTR2 gene described herein). Hybridization probes are oligonucleotides which bind in a base-specific manner to a complementary strand of nucleic acid. Suitable probes include polypeptide nucleic acids, as described in (Nielsen *et al.*, *Science* 254, 1497-1500 (1991)).

Such nucleic acid molecules can be detected and/or isolated by specific hybridization (e.g., under high stringency conditions). "Stringency conditions" for hybridization is a term of art which refers to the incubation and wash conditions, e.g., conditions of temperature and buffer concentration, which permit hybridization of a particular nucleic acid to a second nucleic acid; the first nucleic acid may be perfectly (i.e., 100%) complementary to the second, or the first and second may share some degree of complementarity which is less than perfect (e.g., 60%, 75%, 85%, 95%). For example, certain high stringency conditions can be used which distinguish perfectly complementary nucleic acids from those of less complementarity.

"High stringency conditions", "moderate stringency conditions" and "low stringency conditions" for nucleic acid hybridizations are explained on pages 2.10.1-2.10.16 and pages 6.3.1-6 in Current Protocols in Molecular Biology (Ausubel, F.M. et al., "Current Protocols in Molecular Biology", John Wiley & Sons, (1998)) the 5 teachings of which are hereby incorporated by reference. The exact conditions which determine the stringency of hybridization depend not only on ionic strength (e.g., 0.2XSSC, 0.1XSSC), temperature (e.g., room temperature, 42°C, 68°C) and the concentration of destabilizing agents such as formamide or denaturing agents such as SDS, but also on factors such as the length of the nucleic acid sequence, base 10 composition, percent mismatch between hybridizing sequences and the frequency of occurrence of subsets of that sequence within other non-identical sequences. Thus, high, moderate or low stringency conditions can be determined empirically. By varying hybridization conditions from a level of stringency at which no hybridization occurs to a level at which hybridization is first observed, conditions which will allow a given sequence to hybridize (e.g., selectively) with the most similar sequences in the sample can be determined.

Exemplary conditions are described in Krause, M.H. and S.A. Aaronson, Methods in Enzymology, 200:546-556 (1991). Also, in, Ausubel, et al., "Current Protocols in Molecular Biology", John Wiley & Sons, (1998), which describes the determination of washing conditions for moderate or low stringency conditions. Washing is the step in which conditions are usually set so as to determine a minimum level of complementarity of the hybrids. Generally, starting from the lowest temperature at which only homologous hybridization occurs, each °C by which the final wash temperature is reduced (holding SSC concentration constant) allows an increase by 1% in the maximum extent of mismatching among the sequences that hybridize. Generally, doubling the concentration of SSC results in an increase in T_m of ~17°C. Using these guidelines, the washing temperature can be determined empirically for high, moderate or low stringency, depending on the level of mismatch sought.

For example, a low stringency wash can comprise washing in a solution containing 0.2XSSC/0.1% SDS for 10 min at room temperature; a moderate

25

30

PCT/US00/23021

stringency wash can comprise washing in a prewarmed solution (42°C) solution containing 0.2XSSC/0.1% SDS for 15 min at 42°C; and a high stringency wash can comprise washing in prewarmed (68°C) solution containing 0.1XSSC/0.1%SDS for 15 min at 68°C. Furthermore, washes can be performed repeatedly or sequentially to obtain a desired result as known in the art. Equivalent conditions can be determined by varying one or more of the parameters given as an example, as known in the art, while maintaining a similar degree of identity or similarity between the target nucleic acid molecule and the primer or probe used.

-7-

Hybridizable nucleic acid molecules are useful as probes and primers, e.g., for diagnostic applications, as described below. As used herein, the term "primer" refers to a single-stranded oligonucleotide which acts as a point of initiation of template-directed DNA synthesis under appropriate conditions (e.g., in the presence of four different nucleoside triphosphates and an agent for polymerization, such as, DNA or RNA polymerase or reverse transcriptase) in an appropriate buffer and at a 15 suitable temperature. The appropriate length of a primer depends on the intended use of the primer, but typically ranges from 15 to 30 nucleotides. Short primer molecules generally require cooler temperatures to form sufficiently stable hybrid complexes with the template. A primer need not reflect the exact sequence of the template, but must be sufficiently complementary to hybridize with a template. The 20 term "primer site" refers to the area of the target DNA to which a primer hybridizes. The term "primer pair" refers to a set of primers including a 5' (upstream) primer that hybridizes with the 5' end of the DNA sequence to be amplified and a 3' (downstream) primer that hybridizes with the complement of the 3' end of the sequence to be amplified.

The invention also pertains to nucleotide sequences which have a substantial identity with the nucleotide sequences described herein; particularly preferred are nucleotide sequences which have at least about 70%, and more preferably at least about 80% identity, and even more preferably at least about 90% identity, with nucleotide sequences described herein. Particularly preferred in this instance are nucleotide sequences encoding hypocretin (orexin) receptor 2.

To determine the percent identity of two nucleotide sequences, the sequences are aligned for optimal comparison purposes (e.g., gaps can be introduced in the sequence of a first nucleotide sequence). The nucleotides at corresponding nucleotide positions are then compared. When a position in the first sequence is occupied by the same nucleotide as the corresponding position in the second sequence, then the molecules are identical at that position. The percent identity between the two sequences is a function of the number of identical positions shared by the sequences (i.e., % identity = # of identical positions/total # of positions x 100).

10 The determination of percent identity between two sequences can be accomplished using a mathematical algorithm. A preferred, non-limiting example of a mathematical algorithm utilized for the comparison of two sequences is the algorithm of Karlin et al. (Proc. Natl. Acad. Sci. USA, 90:5873-5877 (1993)). Such an algorithm is incorporated into the NBLAST program which can be used to identify sequences having the desired identity to nucleotide sequences of the invention. To obtain gapped alignments for comparison purposes, Gapped BLAST can be utilized as described in Altschul et al. (Nucleic Acids Res, 25:3389-3402 (1997)). When utilizing BLAST and Gapped BLAST programs, the default parameters of the respective programs (e.g., NBLAST) can be used. See http://www.ncbi.nlm.nih.gov. In one embodiment, parameters for sequence 20 comparison can be set at W=12. Parameters can also be varied (e.g., W=5 or W=20). The value "W" determines how many continuous nucleotides must be identical for the program to identify two sequences as containing regions of identity.

The invention also provides expression vectors containing a nucleic acid

comprising the HCRTR2 gene, operatively linked to at least one regulatory
sequence. Many such vectors are commercially available, and other suitable vectors
can be readily prepared by the skilled artisan. "Operatively linked" is intended to
mean that the nucleic acid sequence is linked to a regulatory sequence in a manner
which allows expression of the nucleic acid sequence. Regulatory sequences are artrecognized and are selected to produce a hypocretin (orexin) receptor 2.
Accordingly, the term "regulatory sequence" includes promoters, enhancers, and

other expression control elements such as those described in Goeddel, Gene Expression Technology: Methods in Enzymology 185, Academic Press, San Diego, CA (1990). For example, the native regulatory sequences or regulatory sequences native to the transformed host cell can be employed. It should be understood that the 5 design of the expression vector may depend on such factors as the choice of the host cell to be transformed and/or the receptor desired to be expressed. For instance, the gene of the present invention can be expressed by ligating the gene into a vector suitable for expression in either prokaryotic cells, eukaryotic cells or both (see, for example, Broach, et al., Experimental Manipulation of Gene Expression, ed. M. Inouye (Academic Press, 1983) p. 83; Molecular Cloning: A Laboratory Manual, 2nd Ed., ed. Sambrook et al. (Cold Spring Harbor Laboratory Press, 1989) Chapters 16 and 17). Typically, expression constructs will contain one or more selectable markers, including, but not limited to, the gene that encodes dihydrofolate reductase and the genes that confer resistance to neomycin, tetracycline, ampicillin, chloramphenicol, kanamycin and streptomycin resistance. Vectors can also include, for example, an autonomously replicating sequence (ARS), expression control sequences, ribosome-binding sites, RNA splice sites, polyadenylation sites, transcriptional terminator sequences, secretion signals and mRNA stabilizing sequences.

20 Prokaryotic and eukaryotic host cells transformed by the described vectors are also provided by this invention. For instance, cells which can be transformed with the vectors of the present invention include, but are not limited to, bacterial cells such as E. coli (e.g., E. coli K12 strains), Streptomyces, Pseudomonas, Serratia marcescens and Salmonella typhimurium, insect cells (baculovirus), including Drosophila, fungal cells, such as yeast cells, plant cells and mammalian cells, such as thymocytes, Chinese hamster ovary cells (CHO), and COS cells. The host cells can be transformed by the described vectors by various methods (e.g., electroporation, transfection using calcium chloride, rubidium chloride, calcium phosphate, DEAE-dextran, or other substances; microprojectile bombardment; 30 lipofection, infection where the vector is an infectious agent such as a retroviral genome, and other methods), depending on the type of cellular host.

The nucleic acid molecules of the present invention can be produced, for example, by replication in a suitable host cell, as described above. Alternatively, the nucleic acid molecules can also be produced by chemical synthesis.

The nucleotide sequences of the nucleic acid molecules described herein

(e.g., a nucleic acid molecule comprising SEQ ID NO:1) can be amplified by methods known in the art. For example, this can be accomplished by e.g., PCR. See generally PCR Technology: Principles and Applications for DNA Amplification (ed. H.A. Erlich, Freeman Press, NY, NY, 1992); PCR Protocols: A Guide to Methods and Applications (eds. Innis, et al., Academic Press, San Diego, CA, 1990); Mattila et al., Nucleic Acids Res. 19, 4967 (1991); Eckert et al., PCR Methods and Applications 1, 17 (1991); PCR (eds. McPherson et al., IRL Press, Oxford); and U.S. Patent 4,683,202.

Other suitable amplification methods include the ligase chain reaction (LCR) (see Wu and Wallace, *Genomics* 4, 560 (1989), Landegren *et al.*, *Science* 241, 1077 (1988), transcription amplification (Kwoh *et al.*, *Proc. Natl. Acad. Sci. USA* 86, 1173 (1989)), and self-sustained sequence replication (Guatelli *et al.*, *Proc. Nat. Acad. Sci. USA*, 87, 1874 (1990)) and nucleic acid based sequence amplification (NASBA). The latter two amplification methods involve isothermal reactions based on isothermal transcription, which produce both single stranded RNA (ssRNA) and double stranded DNA (dsDNA) as the amplification products in a ratio of about 30 or 100 to 1, respectively.

The amplified DNA can be radiolabeled and used as a probe for screening a library or other suitable vector to identify homologous nucleotide sequences. Corresponding clones can be isolated, DNA can be obtained following *in vivo*25 excision, and the cloned insert can be sequenced in either or both orientations by art recognized methods, to identify the correct reading frame encoding a protein of the appropriate molecular weight. For example, the direct analysis of the nucleotide sequence of homologous nucleic acid molecules of the present invention can be accomplished using either the dideoxy chain termination method or the Maxam
Gilbert method (see Sambrook *et al.*, *Molecular Cloning, A Laboratory Manual* (2nd Ed., CSHP, New York 1989); Zyskind *et al.*, *Recombinant DNA Laboratory*

-11-

Manual, (Acad. Press, 1988)). Using these or similar methods, the protein(s) and the DNA encoding the protein can be isolated, sequenced and further characterized.

METHODS OF DIAGNOSIS

The nucleic acids and the proteins described above can be used to detect, in an individual, a mutation in the HCRTR2 gene that is associated with narcolepsy. In 5 one embodiment of the invention, diagnosis of narcolepsy is made by detecting a mutation in the HCRTR2 gene. The mutation can be the insertion or deletion of a single nucleotide, or of more than one nucleotide, resulting in a frame shift mutation; the change of at least one nucleotide, resulting in a change in the encoded amino acid; the change of at least one nucleotide, resulting in the generation of a premature 10 stop codon; the deletion of several nucleotides, resulting in a deletion of one or more amino acids encoded by the nucleotides; the insertion of one or several nucleotides, such as by unequal recombination or gene conversion, resulting in an interruption of the coding sequence of the gene; duplication of all or a part of the gene; transposition of all or a part of the gene; or rearrangement of all or a part of the gene. 15 More than one such mutation may be present in a single gene. Such sequence changes cause a mutation in the receptor encoded by the HCRTR2 gene. For example, if the mutation is a frame shift mutation, the frame shift can result in a change in the encoded amino acids, and/or can result in the generation of a premature stop codon, causing generation of a truncated receptor. Alternatively, a mutation associated with narcolepsy can be a synonymous mutation in one or more nucleotides (i.e., a mutation that does not result in a change in the receptor encoded by the HCRTR2 gene, such as a mutation in an intron or an untranslated portion of the gene). Such a polymorphism may alter splicing sites, affect the stability or 25 transport of mRNA, or otherwise affect the transcription or translation of the gene. A HCRTR2 gene that has any of the mutations described above is referred to herein as a "mutant gene." It is likely that a mutation in the HCRTR2 gene is associated with narcolepsy in humans because of the association between a mutation in the HCRTR2 gene and narcolepsy in dogs (Lin, L. et al., Cell 98:365-376 (1999), the 30 entire teachings of which are incorporated herein by reference). In a preferred

5

30

embodiment, the mutation in the HCRTR2 gene is to a deletion mutation, for example, a deletion that corresponds to the deletions found in the hypocretin (orexin) receptor 2 in narcoleptic dogs as described by Lin et al., supra (e.g., a deletion of one or more exons, such as a deletion of the fourth exon, that can be caused by insertion of one or more nucleotides upstream of the splice site of the exon, or a deletion of exon 6, that can be caused by a G to A transition in the splice junction consensus sequence). In another preferred embodiment, the mutation in the HCRTR2 gene is mutation that effects a "knockout" of the entire gene, such as deletion of the first exon as described by Chemelli, R.M. et al, (Cell 98:437-451 (1999), the entire teachings of which are incorporated herein). In a third preferred embodiment, the mutation in the HCRTR2 gene is a mutation in an intron, that affects splicing (joining of exons) during translation of the HCRTR2 gene.

In a first method of diagnosing narcolepsy, hybridization methods, such as Southern analysis, are used (see Current Protocols in Molecular Biology, Ausubel, F. et al., eds., John Wiley & Sons, including all supplements through 1999). For example, a test sample of genomic DNA, RNA, or cDNA, is obtained from an individual suspected of having (or carrying a defect for) narcolepsy (the "test individual"). The individual can be an adult, child, or fetus. The test sample can be from any source which contains genomic DNA, such as a blood sample, sample of amniotic fluid, sample of cerebrospinal fluid, or tissue sample from skin, muscle, 20 placenta, gastrointestinal tract or other organs. A test sample of DNA from fetal cells or tissue can be obtained by appropriate methods, such as by amniocentesis or chorionic villus sampling. The DNA, RNA, or cDNA sample is then examined to determine whether a mutation in the HCRTR2 gene is present. The presence of the mutation can be indicated by hybridization of the gene in the test sample to a nucleic acid probe. A "nucleic acid probe", as used herein, can be a DNA probe or an RNA probe; the nucleic acid probe contains at least one mutation in the HCRTR2 gene. The probe can be one of the nucleic acid molecules described above (e.g., the gene, a vector comprising the gene, etc.)

To diagnose narcolepsy by hybridization, a hybridization sample is formed by contacting the test sample containing a HCRTR2 gene, with at least one nucleic

15

acid probe. The hybridization sample is maintained under conditions which are sufficient to allow specific hybridization of the nucleic acid probe to the HCRTR2 gene. "Specific hybridization", as used herein, indicates exact hybridization (e.g., with no mismatches). Specific hybridization can be performed under high stringency conditions or moderate stringency conditions, for example, as described above. In a particularly preferred embodiment, the hybridization conditions for specific hybridization are high stringency.

Specific hybridization, if present, is then detected using standard methods. If specific hybridization occurs between the nucleic acid probe and the HCRTR2 gene in the test sample, then the HCRTR2 gene has the mutation that is present in the nucleic acid probe. More than one nucleic acid probe can also be used concurrently in this method. Specific hybridization of any one of the nucleic acid probes is indicative of a mutation in the HCRTR2 gene, and is therefore diagnostic for narcolepsy.

In another hybridization method, Northern analysis (see Current Protocols in Molecular Biology, Ausubel, F. et al., eds., John Wiley & Sons, supra) is used to identify the presence of a mutation associated with narcolepsy. For Northern analysis, a test sample of RNA is obtained from the individual by appropriate means. Specific hybridization of a nucleic acid probe, as described above, to RNA from the individual is indicative of a mutation in the HCRTR2 gene, and is therefore diagnostic for narcolepsy

For representative examples of use of nucleic acid probes, see, for example, U.S. Patents No. 5,288,611 and 4,851,330. Alternatively, a peptide nucleic acid (PNA) probe can be used instead of a nucleic acid probe in the hybridization

25 methods described above. PNA is a DNA mimic having a peptide-like, inorganic backbone, such as N-(2-aminoethyl)glycine units, with an organic base (A, G, C, T or U) attached to the glycine nitrogen via a methylene carbonyl linker (see, for example, Nielsen, P.E. et al., Bioconjugate Chemistry, 1994, 5, American Chemical Society, p. 1 (1994). The PNA probe can be designed to specifically hybridize to a gene having a polymorphism associated with autoimmune disease. Hybridization of the PNA probe to the HCRTR2 gene is diagnostic for narcolepsy..

-14-

In another method of the invention, mutation analysis by restriction digestion can be used to detect mutant genes, or genes containing polymorphisms, if the mutation or polymorphism in the gene results in the creation or elimination of a restriction site. A test sample containing genomic DNA is obtained from the individual. Polymerase chain reaction (PCR) can be used to amplify the HCRTR2 gene (and, if necessary, the flanking sequences) in the test sample of genomic DNA from the test individual. RFLP analysis is conducted as described (see Current Protocols in Molecular Biology, supra). The digestion pattern of the relevant DNA fragment indicates the presence or absence of the mutation in the HCRTR2 gene, and therefore indicates the presence or absence of narcolepsy.

Sequence analysis can also be used to detect specific mutations in the HCRTR2 gene. A test sample of DNA is obtained from the test individual. PCR can be used to amplify the gene, and/or its flanking sequences. The sequence of the HCRTR2 gene, or a fragment of the gene is determined, using standard methods. The sequence of the gene (or gene fragment) is compared with the nucleic acid sequence of the gene, as described above. The presence of a mutation in the HCRTR2 gene indicates that the individual has narcolepsy.

10

15

20

25

Allele-specific oligonucleotides can also be used to detect the presence of a mutation in the HCRTR2 gene, through the use of dot-blot hybridization of amplified proteins with allele-specific oligonucleotide (ASO) probes (see, for example, Saiki, R. et al., (1986), Nature (London) 324:163-166). An "allele-specific oligonucleotide" (also referred to herein as an "allele-specific oligonucleotide probe") is an oligonucleotide of approximately 10-50 base pairs, preferably approximately 15-30 base pairs, that specifically hybridizes to the HCRTR2 gene, and that contains a mutation associated with narcolepsy. An allele-specific oligonucleotide probe that is specific for particular mutation in the HCRTR2 gene can be prepared, using standard methods (see Current Protocols in Molecular Biology, supra). To identify mutations in the gene that are associated with narcolepsy, a test sample of DNA is obtained from the individual. PCR can be used 30 to amplify all or a fragment of the HCRTR2 gene, and its flanking sequences. The DNA containing the amplified HCRTR2 gene (or fragment of the gene) is dotblotted, using standard methods (see Current Protocols in Molecular Biology, supra), and the blot is contacted with the oligonucleotide probe. The presence of specific hybridization of the probe to the amplified HCRTR2 gene is then detected. Specific hybridization of an allele-specific oligonucleotide probe to DNA from the individual is indicative of a mutation in the HCRTR2 gene, and is therefore indicative of narcolepsy.

Other methods of nucleic acid analysis can be used to detect mutations in the HCRTR2 gene, for the diagnosis of narcolepsy. Representative methods include direct manual sequencing; automated fluorescent sequencing; single-stranded conformation polymorphism assays (SSCA); clamped denaturing gel electrophoresis (CDGE) heteoduplex analysis; chemical mismatch cleavage (CMC); RNase protection assays; use of proteins which recognize nucleotide mismatches, such as *E. coli* mutS protein; allele-specific PCR, and other methods.

PHARMACEUTICAL COMPOSITIONS

The present invention also pertains to pharmaceutical compositions comprising nucleic acids described herein, particularly nucleic acids containing the HCRTR2 gene described herein. For instance, a nucleotide or nucleic acid construct (vector) comprising a nucleotide of the present invention can be formulated with a physiologically acceptable carrier or excipient to prepare a pharmaceutical composition. The carrier and composition can be sterile. The formulation should suit the mode of administration.

Suitable pharmaceutically acceptable carriers include but are not limited to water, salt solutions (e.g., NaCl), saline, buffered saline, alcohols, glycerol, ethanol, gum arabic, vegetable oils, benzyl alcohols, polyethylene glycols, gelatin, carbohydrates such as lactose, amylose or starch, dextrose, magnesium stearate, talc, silicic acid, viscous paraffin, perfume oil, fatty acid esters, hydroxymethylcellulose, polyvinyl pyrolidone, etc., as well as combinations thereof. The pharmaceutical preparations can, if desired, be mixed with auxiliary agents, e.g., lubricants, preservatives, stabilizers, wetting agents, emulsifiers, salts for influencing osmotic

-16-

pressure, buffers, coloring, flavoring and/or aromatic substances and the like which do not deleteriously react with the active compounds.

The composition, if desired, can also contain minor amounts of wetting or emulsifying agents, or pH buffering agents. The composition can be a liquid solution, suspension, emulsion, tablet, pill, capsule, sustained release formulation, or powder. The composition can be formulated as a suppository, with traditional binders and carriers such as triglycerides. Oral formulation can include standard carriers such as pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, polyvinyl pyrollidone, sodium saccharine, cellulose, magnesium carbonate, etc.

10

15

25

30

Methods of introduction of these compositions include, but are not limited to, intradermal, intramuscular, intraperitoneal, intraocular, intravenous, subcutaneous, oral and intranasal. Other suitable methods of introduction can also include gene therapy (as described below), rechargeable or biodegradable devices, particle acceleration devises ("gene guns") and slow release polymeric devices. The pharmaceutical compositions of this invention can also be administered as part of a combinatorial therapy with other agents.

The composition can be formulated in accordance with the routine procedures as a pharmaceutical composition adapted for administration to human 20 beings. For example, compositions for intravenous administration typically are solutions in sterile isotonic aqueous buffer. Where necessary, the composition may also include a solubilizing agent and a local anesthetic to ease pain at the site of the injection. Generally, the ingredients are supplied either separately or mixed together in unit dosage form, for example, as a dry lyophilized powder or water free concentrate in a hermetically sealed container such as an ampoule or sachette indicating the quantity of active agent. Where the composition is to be administered by infusion, it can be dispensed with an infusion bottle containing sterile pharmaceutical grade water, saline or dextrose/water. Where the composition is administered by injection, an ampoule of sterile water for injection or saline can be provided so that the ingredients may be mixed prior to administration.

-17-

For topical application, nonsprayable forms, viscous to semi-solid or solid forms comprising a carrier compatible with topical application and having a dynamic viscosity preferably greater than water, can be employed. Suitable formulations include but are not limited to solutions, suspensions, emulsions, creams, ointments, powders, enemas, lotions, sols, liniments, salves, aerosols, etc., which are, if desired, sterilized or mixed with auxiliary agents, e.g., preservatives, stabilizers, wetting agents, buffers or salts for influencing osmotic pressure, etc. The agent may be incorporated into a cosmetic formulation. For topical application, also suitable are sprayable aerosol preparations wherein the active ingredient, preferably in combination with a solid or liquid inert carrier material, is packaged in a squeeze bottle or in admixture with a pressurized volatile, normally gaseous propellant, e.g., pressurized air.

Agents described herein can be formulated as neutral or salt forms.

Pharmaceutically acceptable salts include those formed with free amino groups such as those derived from hydrochloric, phosphoric, acetic, oxalic, tartaric acids, etc., and those formed with free carboxyl groups such as those derived from sodium, potassium, ammonium, calcium, ferric hydroxides, isopropylamine, triethylamine, 2-ethylamino ethanol, histidine, procaine, etc.

The agents are administered in a therapeutically effective amount. The amount of agents which will be therapeutically effective in the treatment of narcolepsy can be determined by standard clinical techniques. In addition, *in vitro* or *in vivo* assays may optionally be employed to help identify optimal dosage ranges. The precise dose to be employed in the formulation will also depend on the route of administration, and the seriousness of the disease or disorder, and should be decided according to the judgment of a practitioner and each patient's circumstances. Effective doses may be extrapolated from dose-response curves derived from *in vitro* or animal model test systems.

20

25

30

The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Optionally associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture,

-18-

use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use of sale for human administration. The pack or kit can be labeled with information regarding mode of administration, sequence of drug administration (e.g., separately, sequentially or concurrently), or the like. The pack or kit may also include means for reminding the patient to take the therapy. The pack or kit can be a single unit dosage of the combination therapy or it can be a plurality of unit dosages. In particular, the agents can be separated, mixed together in any combination, present in a single vial or tablet. Agents assembled in a blister pack or other dispensing means is preferred. For the purpose of this invention, unit dosage is intended to mean a dosage that is dependent on the individual pharmacodynamics of each agent and administered in FDA approved dosages in standard time courses.

METHODS OF THERAPY

The present invention also pertains to methods of therapy for narcolepsy, 15 utilizing the pharmaceutical compositions comprising nucleic acids, as described herein. The therapy is designed to replace/supplement activity of the hypocretin(orexin) receptor 2 in an individual, such as by administering a nucleic acid comprising the HCRTR2 gene or a derivative or active fragment thereof. In one embodiment of the invention, a nucleic acid of the invention is used in the treatment 20 of narcolepsy. The term, "treatment" as used herein, refers not only to ameliorating symptoms associated with the disease, but also preventing or delaying the onset of the disease, and also lessening the severity or frequency of symptoms of the disease. In this embodiment, a nucleic acid of the invention (e.g., the HCRTR2 gene (SEQ ID NO:1)) can be used, either alone or in a pharmaceutical composition as described above. For example, the HCRTR2 gene, either by itself or included within a vector, can be introduced into cells (either in vitro or in vivo) such that the cells produce native HCRTR2 receptor. If necessary, cells that have been transformed with the gene or can be introduced (or re-introduced) into an individual affected with the disease. Thus, cells which, in nature, lack native HCRTR2 expression and activity, 30 or have mutant HCRTR2 expression and activity, can be engineered to express

-19-

HCRTR2 receptors (or, for example, an active fragment of the HCRTR2 receptor). In a preferred embodiment, nucleic acid comprising the HCRTR2 gene, can be introduced into an expression vector, such as a viral vector, and the vector can be introduced into appropriate cells which lack native HCRTR2 expression in an animal. In such methods, a cell population can be engineered to inducibly or constitutively express active HCRTR2 receptor. Other gene transfer systems, including viral and nonviral transfer systems, can be used. Alternatively, nonviral gene transfer methods, such as calcium phosphate coprecipitation, mechanical techniques (e.g., microinjection); membrane fusion-mediated transfer via liposomes; or direct DNA uptake, can also be used.

10

The nucleic acids and/or vectors are administered in a therapeutically effective amount (i.e., an amount that is sufficient to treat the disease, such as by ameliorating symptoms associated with the disease, preventing or delaying the onset of the disease, and/or also lessening the severity or frequency of symptoms of the disease). The amount which will be therapeutically effective in the treatment of a particular disorder or condition will depend on the nature of the disorder or condition, and can be determined by standard clinical techniques. In addition, in vitro or in vivo assays may optionally be employed to help identify optimal dosage ranges. The precise dose to be employed in the formulation will also depend on the route of administration, and the seriousness of the disease or disorder, and should be decided according to the judgment of a practitioner and each patient's circumstances. Effective doses may be extrapolated from dose-response curves derived from in vitro or animal model test systems.

The following Examples are offered for the purpose of illustrating the present invention and are not to be construed to limit the scope of this invention. The teachings of all references cited herein are hereby incorporated herein by reference.

-20-

EXAMPLES

EXAMPLE 1 Identification of the Human Narcolepsy Gene

A human BAC library (RPCI11 human male BAC library; see Osoegawa, K. et al., Genomics 52:1-8 (1998)) was used. Twenty primers, designed from the mRNA sequence of the HCRTR2 receptor, were employed to identify clones of interest. They are set forth in Table 1.

TABLE 1 Primers Used for Hybridization

	#	Name	Primer Sequence	SEQ ID NO:
5	1	HCRTR2-1-F	TACTACTAGGCCACGCG	3
	2	HCRTR2-1-R	ACACCAGGAGGAGAAAGCTAC	4
	3	HCRTR2-2-F	ATCGCCTGTAAAGACAGCAAAG	5
·	4	HCRTR2-2-R	AAAGTTACTGAGCCAATGCCTC	6
	5	HCRTR2-3-F	GAGAGGAGCTTGCAGCATTG	7
	6	HCRTR2-3-R	AGGAATTCCTCGTCGTCATAGT	8
	7	HCRTR2-4-F	GAAGAACCACCACATGAGGAC	9
10	8	HCRTR2-4-R	ATCACTTTGCAAAGGGACTGTC	10
	9	HCRTR2-5-F	GTATGCAATCTGTCACCCTTTG	11
	10	HCRTR2-5-R	AATGCAGGAGACAATCCAGATG	12
15	11	HCRTR2-6-F	CAGGCTTAGCCAATAAAACCAC	13
	12	HCRTR2-6-R	GATAAGCCAACACCATGAGACA	14
	13	HCRTR2-7-F	ACAGATCCCTGGAACATCATCT	15
	14	HCRTR2-7-R	CTCGGATCTGCTTTATTTCAGC	16
	15	HCRTR2-8-F	CCAATTAGCATCCTCAATGTGC	17
	16	HCRTR2-8-R	GTGTGAAAAGGTAAACCAGGCA	18
20	17	HCRTR2-9-F	CTCAGTGGAAAATTTCGAGAGG	19
	18	HCRTR2-9-R	GTTGCTGATTTGAGTGGTCAAG	20
	19	HCRTR2-10-F	CTTTCTGAGCAAGTTGTGCTCA	21
	20	HCRTR2-10-R	TACCAGTTTTGAAGTGGTCCTG	22

Initial Study with Large Membranes

Four out of 5 membranes having the whole BAC library, containing a total of approximately 160,000 BAC clones representing an approximately 10-fold coverage of the human genome, were used in hybridization studies with these primers.

Hybridization was performed with a pool of all 20 primers described in Table 1.

5' End Labeling for Big Membranes

Oligonucleotides were labeled at the 5' end before hybridization, using fresh (less than one month old) [γ^{32} P]ATP (6000 Ci/mmole; 10 μ Ci/ μ l). The following protocol is adjusted for 4 membranes in 2 bottles, containing 2 membranes/30 ml of rapid hyb. Each. Briefly, a labeling mixture was made of DNA (8 pmol/ μ l) (10.0 μ l of the primer pool), 10X buffer (12.0 μ l), T4 PNK (10 ν l) (6.0 μ l), [γ^{32} P]ATP (30.0 μ l, or 600 μ Ci), and water (62.0 μ l) for a final volume of 120 μ l. 20 μ l of labeling mixture was used per 10 ml rapid hybridization reaction. Incubation of the labeling mixture was for 2 hours at 37°C, followed by transfer to ice, spinning down, and mixing with the rapid hybridization solution. The membranes were prehybridized at 42°C before the labeling mix was added. Sixty μ l of the labeling mix was added to each of 2 big bottles containing 2 membranes and 30 ml of rapid hybridization solution.

Hybridization and Washing

15

The membranes were hybridized at 42°C overnight. After overnight, membranes were washed with 6x SSC, 0.1% SDS at room temperature; washed with 6x SSC, 0.1% SDS at 55°C in a shaking waterbath, repeated until the radioactivity of membranes was lower than 6k using 1x sensitivity; and washed with 6x SSC to remove the SDS. The washed membranes were put in a cassette for overnight exposure at -80°C with a MR single emulsion film. Positive clones were identified and gridded on small membranes.

Study of Positive Clones with Small Membranes

After growing the positively-identified clones on several small membranes (to get several copies of membranes containing the same clones), and washing the membranes, hybridization was performed using pairs of primers, instead of a total pool of primers as before. The total number of hybridizations was ten, using different primers against identical copies of membranes containing all positive clones from the first hybridization. The primer pairs are set forth in Table 2; primer numbers indicate the primers shown in Table 1.

TABLE 2 Primer Pairs Used for Hybridization

	Reaction number	Primers Used	
	1	1 and 2	
	2	3 and 4	
5	3	5 and 6	
	4	7 and 8	
	5	9 and 10	
	6	11 and 12	
	7	13 and 14	
10	8	15 and 16	
	9	17 and 18	
	10	18 and 19	

5' End Labeling for Small Membranes

Oligonucleotides were labeled at the 5' end before hybridization, using fresh

[γ³²P]ATP (5000 Ci/mmole; 10 μCi/μl). Briefly, a labeling mixture was made of DNA (8 pmol/μl) (1.5 μl), 10X buffer (2.0 μl), T4 PNK (10 u/μl) (1.0 μl),

[γ³²P]ATP (3.0 μl), and water (12.5 μl) for a final volume of 20 μl. Incubation of the labeling mixture was for 2.5 hours at 37°C, followed by transfer to ice, spinning down, and mixing with the rapid hybridization solution. Membranes were prewetted in 6X SSC, rolled in a pipette, and excess liquid drained prior to placing the membrane in the tube. Fifty ml Falcon (polypropylene) tubes were used as container for the hybridization. The membranes were prehybridized at 42°C before 20 μl of labeling mix was added to each tube.

Hybridization and Washing

The membranes were hybridized at 42°C overnight. After overnight, membranes were washed as described above. Four clones which were positive for primers designed using the 5' and 3' end of the mRNA were identified. Clone 403B19 was used to characterize the gene.

Sequencing of Narcolepsy Gene in Clone 403B19

Shotgun sequencing was used to obtain the gene sequence.

Preparation of DNA Samples

BAC DNA was isolated using the Plasmix kit from TALENT-VH Bio 5 Limited. Thirty µg of isolated DNA was fragmented by nebulization: a nebulizer (IPI Medical Products, Inc., no. 4207) was modified by removing the plastic cylinder drip ring, cutting off the outer rim of the cylinder, inverting it and placing it back into the nebulizer; the large hole in the top cover (where the mouth piece was attached) was sealed with a plastic stopper; the small hole was connected to a 1/4 inch length of Tycon tubing (connected to a compressed air source). A DNA sample was prepared containing 30 μg DNA, 10 X TM buffer (200 μl), sterile glycerol (1 ml), and sterile dd water (q.s.) for a total volume of 2 ml. The DNA sample was nebulized in an ice-water bath for 2 minutes and 40 seconds (pressure bar reading 15 0.5). The sample was then briefly centrifuged at 2500 rpm to collect the DNA; the entire unit was placed in the rotor bucked of a table top centrifuge (Beckman GPR tabletop centrifuge) fitted with pieces of Styrofoam to cushion the nebulizer. The sample was then distributed into four 1.5 ml microcentrifuge tubes and ethanol precipitated. The Dried DNA pellet was resuspended in 35 μl of 1X TM buffer prior to proceeding with fragment end-repair.

Fragment End Repair, Size Selection and Phosphorylation

The DNA was resuspended in 27 μl of 1X TM buffer. The following materials were added: 10 X kinase buffer (5 μl), 10 mM rATP (5 μl), 0.25 mM

25 dNTPs (7 μl), T4 polynucleotide kinase (1 μl (3 U/μl)), Klenow DNA polymerase (2 μl (5 U/μl)), T4 DNA polymerase (1 μl (3 U/μl)), for a total volume of 48 μl. The mixture was incubated at 37°C for 30 minutes, and then 5 μl of agarose gel loading dye was added. The mixture was then applied to separate wells of a 1% low melting temperature agarose gel and electrophoresed for 30-60 minutes at 100-120 mA. The DNA was then eluted from each sample lane, extracted from the agarose

-25-

using Ultrafree-DA columns (Millipore) and then cleaned with Microcon-100 columns (Amicon), precipitated in ethanol, and resuspended in 10 μ l of 10:0.1 TE buffer.

Ligation

EcoRV-linearized, CIAP-dephosphorylated Bluescript vector was used as a cloning vector. The following reagents were combined in a microcentrifuge tube, and incubated overnight at 4°C: DNA fragments (100-1000 ng), cloning vector (2 μl (10 ng/μl)), 10X ligation buffer (1 μl), T4 DNA ligase (NEB 202L) (1 μl (400 U/μl)), sterile dd water (q.s.), for a total of 10 μl.

10 Transformation of Ligated Products

The ligation products were diluted 1:5 with dd water and used to transform electrocompetent TOP 10F cells (Invitrogen) using GenePulser II (Biorad; voltage, 2.5 W, resistance 100 ohm). Transformants were plated on LB plates with 50 µl of 4% X-GAL and 50 µl of 4% IPTG, and ampicillin. Transformants were grown overnight at 37°C, white colonies were picked, grown in a culture of 3 ml LB liquid media plus 200 µg/µl ampicillin for 16-20 hours with shaking. DNA was isolated from the liquid cultures using Autogen 740 Automatic Plasmid Isolation System.

Cycle Sequencing of Isolated Plasmid DNA

Isolated plasmids were then sequenced using the M13 primers: M13-forward (SEQ ID NO:23) TGTAAAACGACGGCCAG; and M13-reverse (SEQ ID NO:24) CAGGAAACAGCTATGAC. For the sequencing reaction, 2.5 µl plasmid template was mixed with 4 µl Big Dye Ready reaction mix (ABI), 1 µl of 8 pM M13 primer, and 2.5 µl dd water. For cycle sequencing, 25 cycles of 96°C for 10 seconds, 50°C for 5 seconds, and 60 °C for 4 minutes were performed, followed by holding at 4°C.

The cycle sequencing reaction products were cleaned by spinning through Sephadex G-50 columns. The eluted cycle sequencing products were then dissolved in 3 µl formamide/dye and 1.5 µl of sample was loaded on ABI 377 automated sequencers. The data was analyzed using Phred and Phrap (Ewing, B. et al., Genome Res. 8:175-

185 (1998); Ewing, B. and Green, P., Genome Res. 8:186-194 (1998)), and viewed in Consed viewer (Gordon, D. et al., Genome Res. 8(3):195-202 (1998)).

Analysis of Gene Structure

The *hcrtr-2* gene maps to chromosome 6p11-q11. A total of 168,575 base pairs of contiguous sequence was generated for 403B19 which contained all of the *hcrtr-2* gene. Comparison of the cDNA sequence of *hcrtr-2* (Accession number GI:6006037) and the genomic sequences generated allowed deduction of the intron/exon organization of the gene. The gene contains 7 exons which cover 108,439 bp. The first 10 Gs in the mRNA sequence for *hcrtr-2* were not found in the genomic sequence. It is likely that these Gs were an artifact.

The splice junctions of the *hcrtr-2* gene are set forth in Table 3, and the intron sizes are set forth in Table 4. Exon sequences are represented in uppercase and introns in lowercase. All splice sites conform to the consensus GT-AG rule. SEQ ID NOs are given in the column immediately following each site.

15 Table 3 Splice Junctions of hcrtr-2

	Splice Donor Site	SEQ ID	Splice Acceptor Site	SEQ ID
Hcrtr-2 exon1-2	TCCTGGgtgagt	25	aattagTTTGTG	26
Hcrtr-2 exon2-3	CTACAGgtaatt	27	ctctagACCGTG	28
Hcrtr-2 exon3-4	GGGGTGgtaagt	29	tcctagGTGAAA	30
Hcrtr-2 exon4-5	CGACAGgtatat	31	tttcagATCCCT	32
Hcrtr-2 exon5-6	AAAGAGgtaaaa	33	ctgcagAGTATT	34
Hcrtr-2 exon6-7	TCAGTGgtgagt	35	tgccagGAAAAT	36

Table 4 Intron Sizes of hcrtr-2

Intron	Nucleotides
Intron 1	73,848
Intron 2	6,322
Intron 3	8,327
Intron 4	13,618
Intron 5	2,730
Intron 6	1,779

The exons do not clearly respect the domain structure of this seven

membrane domain G protein linked receptor. Five of the transmembrane regions are
by themselves within one exon, two of the transmebrane segments are broken up by
introns, and two transmembrane segments fall within the same exon. A survey done
one year ago on mammalian G-protein coupled receptors (GPCRs) sequences in
GenBank revealed that over 90% of GPCRs genes were intronless in their open

reading frame (ORF) (Gentles, A.J. and Karlin, S., Trends Genet. 15:47-49 (1999)).
Comparison of the intron/exon boundaries of hcrtr-2 and the genes coding for their
most related GPCRs based on sequence similarity showed that the location of the
intron/exons boundaries with respect to the transmembrane domains is only partially
conserved among the receptors (Sakurai, T. et al., Cell 92:573-585 (1998)).

20 Computer analysis of sequence data

Analysis of the genomic sequence of hcrtr-2 using the program RepeatMasker (http://ftp.genome.washington.edu/cgi-bin/RepeatMasker) showed that the sequence containing the hcrtr-2 genomic sequence is 38.27% repeat sequences and the GC content is 35.3%.

The sequences of the genes were analyzed using the program GeneMiner (Óskarsson and Pálsson, unpublished), which combines the results of 5 exon prediction programs; FGENE (Solovyev, V. and Salamov, A., Ismb 5:294-302 (1997)), Genscan (Burge, C. and Karlin, S., J. Mol. Biol. 268:78-94 (1997)),

HMMgene (Krogh, A., Ismb 5:179-186 (1997)), MZEF (Zhang, M.Q., Proc. Natl. Acad. Sci. USA 94:565-8 (1997)) and Xpound (Thomas, A. and Skolnick, M.H., IMA J. Math Appl. Med. Biol. 11:149-160 (1994)). For hcrtr-2, 3 out of 5 programs predicted the 3' end of exon 1, only one program predicted the 7th exon and for the internal exons, there were at least two programs that predicted each of them exactly or in part.

The promoter sequences of the genes have not yet been characterized. The Promoter Prediction by Neural Network

(http://www.fruitfly.org/seq_tools/promoter.html) predicted promoters that are at least

10 140 bp upstream of the 5'UTR of hcrtr-2, indicating that either a part of the 5'UTR is missing in the published mRNA sequence or the real promoters are not detected by the program.

Analysis of Population for Polymorphisms

in nucleic acid samples from 47 patients and 75 control individuals. The patient population consisted of patients of Icelandic and US origin. The control population consisted of Icelandic controls, CEPH (Centre d'Etude du Polymorphisme Humain) individuals from Utah and France, and US samples of various ethnic origins. The African-American/Caucasian ratios were similar between patients and controls. All narcoleptic subjects complained of excessive daytime sleepiness (EDS). Approximately 66% of the patients had cataplexy, 24% did not and 10% did not have attainable records of cataplexy status. Narcoleptic subjects without cataplexy had Multiple Sleep Latency Tests showing mean sleep latencies of less than 10 minutes and REM sleep in at least 2 naps. Subjects did not take any drugs affecting sleep for at least 10 days before their sleep studies.

To analyze the nucleic acids, DNA from patient and control blood samples were isolated by the method of Kunkel (Kunkel, L.M. et al., Proc. Natl. Acad. Sci. USA 74:1245-9 (1977)). Briefly, white blood cells were lysed in a sucrose lysis buffer, and proteinase K treated; the DNA was then extracted using phenol-chloroform/isoamylalcohol and then ethanol precipitated. Patient samples that were

received in the form of nuclei pelleted through sucrose buffer were resusupended in lysis buffer (100 mM NaCl2; 10 mM TrisHCl, pH 8; 25 mM EDTA pH 8; 0.5% sodium dodecyl sulfate; 0.1 mg/ml proteinase K) at 55°C for 4-6 hours followed by classical phenol-chloroform extraction and ethanol precipitation (Sambrook, J. et al., Molecular Cloning, A Laboratory Manual (1989)). Samples were incubated at 55°C after isolation for the inactivation of DNAse to prevent the degradation of DNA. Concentration of the isolated DNA was determined by spectrophotometric analysis at 260 nm (Sambrook et al., using GeneQuant (PharmaciaBiotech), and samples

Primers were designed from intronic sequences flanking the exons of the hypocretin receptor-2 (*hcrtr-2*), using either primer design programs available at primer3 at the Whitehead Institute (http://www-genome.wi.mit.edu/cgi-bin/primer3.cgi) or primers for the worldwide web (http://williamstone.com/primers/javascript/). The primers are shown in Table 5.

diluted with sterile distilled water to a 20 ng/µl working solution.

Table 5 Primers Used to Amplify Nucleic Acid Fragments for Analysis of hcrtr-2 Gene

	EX-	#	Primer Sequence	Sense/	External/	SEQ
	ON	_		Antisense	Nested	ID.
5	1	1	TTTCTTCAGCTTCAGCTCTCCCTCAGC	S	Е	37
	1	2	TTCAGCTCCGAAGCAGATGACCAGTTG	A	Е	38
	1	3	TTCAGCTTCAGCTCTCCCTCAGCGAGG	S	N	39
	1	4	CGAAGCAGATGACCAGTTGCGACAAGG	A	N	40
	1	5	CTTTCCCACCGCAAATCACCAGTGCTC	s	Е	41
10	1	6	ATTITATTAGAAAACCCCATCCGAGAG	A	E	42
	1	7	TTCCCACCGCAAATCACCAGTGCTC	s	N	43
	1	8	TATTAGAAAACCCCATCCGAGAGCAG	A	N	44
	2	9	GCATGTACTTAGCATTCACACAGATTG	S	E	45
	2	10	TCTAATGATGATTTGGCAGTTCATTGC	A	E	46
15	2	11	TAGCATTCACACAGATTGACAGATTCA	s	N	47
	2	12	CAGTTTGTCAATGCCTTAGGCAAATAT	A	N	48
	3	13	TTTGGCAGCTTTGAATTTGCTTATATG	S	E	49
	3	14	GCTCTTGCAAAACTGTATTCACAAATG	Α	E	50
	3	15	CAGCITIGAATTTGCTTATATGTTGTG	S	N	51
20	3	16	TTGCAAAACTGTATTCACAAATGTCAA	Α	N	52
	4	17	TCCCCTTTGCATACATAATATGACAATG	S	Е	53
	4	18	AAAAAGCACAGACAAAATATTTGGAAGG	Α	Е	54
	4	19	ATGCACTTTGAAGAAAAGCATTGACATG	s	N	55
	4	20	AAGCACAGACAAAATATTTGGAAGGAAT	A	N	56
25	5	21	CTCAGGCGTCTGGAAGCCTTTCCTTAC	S	Е	57
	5	22	TTAAAGGCTGTTCGCCTTACCTGCTGG	Α	Е	58
	5	23	GGCGTCTGGAAGCCTTTCCTTACTGTG	s	N	59
	5	24	CTGAGTCATCTGGCCTGACAAGGTATC	A	N	60
	6	25	GGGTCAGAAACCAATCTGTGGTCAATTC	s	E	61
30	6	26	AGTTGAAGAGTGTTCATTGATTCCTCATCC	Α	E	62
	6	27	AGAAACCAATCTGTGGTCAATTCCTGCAAC	S	N	63

-31-

EX-	#	Primer Sequence	Sense/	External/	SEQ
ON			Antisense	Nested	ID.
6	28	TGAAGAGTGTTCATTGATTCCTCATCCTTG	A	N	64
7	29	GAGTCTACCAAGCTTCCAATAAACTCA	S	E	65
7	30	GGATAGTTTTACTCAGGTATCCTTGTCA	A	E	66
7	31	CAAATCAGCAACTTTGATAACATAT	S	N	67
7	32	GTATCCTTGTCATATGAATAAATATTCTAC	A	N	68
7	33	CACTCAAATCAGCAACTTTGATAAC	S	E	69
7	34	GTGAGAGATTAAAATAACAAGGGAT	A	E	70
7	35	CAAATCAGCAACTTTGATAACATAT	s	N	71
7	36	TGTTTAAACATTTAATTGACACACA	A	N	72
7	37	TTCATATGACAAGGATACCTGAGTAAA	S	Е	73
7	38	GTGAAATAGCCTGAAATAAGCTCAA	A	Е	74

5

10

PCR reactions were done in 20 µl reactions using 40 ng genomic DNA, 0.2 mM solution of the four dNTPs, 0.35 µM of each primer (TAGCopenhagen), 2.5 mM MgCl2 (Perkin Elmer), 1x PCR Buffer (Perkin Elmer) and 0.5 U Amplitaq gold 15 (Perkin Elmer). The primers were used to amplify the fragments by PCR cycling at 95°C for 12 min and subsequently 30 cycles of 95°C for 30 sec, 55-62°C for 30 sec and 72°C for 1 min. The PCR products were prepared for cycle sequencing by incubation with Shrimp alkaline phosphatase (Amersham) and exonuclease I (Amersham) at 37°C for 15 min. After the inactivation of the enzymes the products were subject to cycle sequencing using BigDye Ready Reaction mix (Perkin Elmer) 20 and subsequently run on ABI Prism 377 Automated DNA sequencers. The raw data were basecalled and sequences assembled using the Phred and Phrap software, respectively (Ewing, B. et al., Genome Res. 8:175-185 (1998); Ewing, B. and Green, P., Genome Res. 8:186-194 (1998)). The Consed viewer was used to analyze the sequences (Gordon, D. et al., Genome Res. 8(3):195-202 (1998)). Expansion of a Tstretch in the 3' untranslated region (UTR) of exon 7 of hcrtr-2 was investigated by amplifying a fragment containing the stretch with a fluorescently labelled primer

pair using the conditions described above. The PCR product was dissolved in formamide/dye solution and run on ABI Prism 377 Automated DNA sequencers as described above. Allele calling was done using TrueAllele and editing was done using DeCODE-GT (Palsson, B. et al., Genome Res. 9:1002-1012 (1999)).

A total of nine single nucleotide polymorphisms were identified, 7 in exons and 2 in an intronic sequence near an exon. The polymporphisms are shown in Table 6. The base number is according to the mRNA sequence (Accession number GI:6006037). For those polymorphisms marked with an asterisk (*), the polymorphism is located 5' of the corresponding exons; the numbers indicate the distance into the introns.

Table 6 Single Nucleotide Polymorphisms in hcrtr-2

5

10

	Location	cDNA base #	Nucleic Acid Change
	Exon 1	352	С-Т
	Exon 1	355	C-A
15	Intron1	-26*	C-A
	Exon 5	1,170	G-A
	Exon 5	1,177	C-A
	Exon 5	1,201	G-A
	Exon 5	1,246	G-A
20	Exon 5	1,266	G-A
	Intron 6	-87*	G-A

While this invention has been particularly shown and described with references to preferred embodiments thereof, it will be understood by those skilled in the art that various changes in form and details may be made therein without departing from the spirit and scope of the invention as defined by the appended claims.

-33-

CLAIMS

What is claimed is:

5

- Isolated nucleic acid molecule comprising the nucleic acid having SEQ ID NO:1.
- A DNA construct comprising the isolated nucleic acid molecule of Claim 1 operatively linked to a regulatory sequence.
- 3. A recombinant host cell comprising the isolated nucleic acid molecule of Claim 1 operatively linked to a regulatory sequence.
- 10 4. A pharmaceutical composition comprising a nucleic acid comprising the isolated nucleic acid molecule of Claim 1.
 - Isolated nucleic acid molecule comprising the nucleic acid having SEQ ID
 NO:1 with one or more of the nucleic acid changes shown in Table 6.
- 6. A method of diagnosing narcolepsy in an individual, comprising detecting a

 mutation in the gene encoding hypocretin (orexin) receptor 2, wherein the

 presence of the mutation in the gene is indicative of narcolepsy.
 - 7. A method of treating narcolepsy in an individual, comprising administering to the individual an isolated nucleic acid of Claim 1 in a therapeutically effective amount.

1/51

```
LOCUS
                                    168,575 bp DNA
                                                          PRI
                                                                 20-OCT-1999
DEFINITION
              Human hypocretin (orexin) receptor 2 (HCRTR2) gene, complete cds.
ACCESSION
NID
VERSION
KEYWORDS
SOURCE
              human.
 ORGANISM
              Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
              Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
              1 (bases 1-168,575)
AUTHORS
 TITLE
              Direct Submission
JOURNAL
                      Submitted (
                                         ) deCode Genetics, Inc., Lynghals 1,
                     IS-110 Reykjavík, Iceland.
FEATURES
                      Location/Qualifiers
       source
                      1.. 168,575
                     /organism="Homo sapiens"
                     /db xref="taxon: 9606"
                     /chromosome="6"
                     /map="6p11-q11"
                     /clone="BAC 403B19"
       gene
                     1..129,305
                     /partial
                     /gene="HCRTR2"
                     /note="OX2R"
                     /db_xref="LocusID:3062"
                     /db_xref="MIM:602393"
                     20,867..21,403
      exon
                     /gene="HCRTR2"
                     /number=2
      CDS
                     join(21,181..21,403, 95,252..95,430, 101,753..101,996, 110,324..110,439,
                     124,058..124,278, 127,009..127,130, 128,910..129,139)
                     /gene="HCRTR2"
                     /note="HCRTR2 exons defined by comparison to mRNA sequence (NM_001526)"
                            /product="HCRTR2/orexin 2 receptor"
                     /db_xref="LocusID:3062"
                     /db_xref="MIM:602393"
                     /protein_id="NP_001517.1"
                     /db_xref="PID:g4557639"
                     /db xref="GI:4557639"
                     translation="MSGTKLEDSPPCRNWSSASELNETQEPFLNPTDYDDEEFLRYLW/
                     REYLHPKEYEWVLIAGYIIVFVVALIGNVLVCVAVWKNHHMRTVTNYFIVNLSLADVL
                     VTITCLPATLVVDITETWFFGQSLCKVIPYLQTVSVSVSVLTLSCIALDRWYAICHPL
                    MFKSTAKRARNSIVIIWIVSCIIMIPQAIVMECSTVFPGLANKTTLFTVCDERWGGEI
                    YPKMYHICFFLVTYMAPLCLMVLAYLQIFRKLWCRQIPGTSSVVQRKWKPLQPVSQPR
                    GPGQPTKSRMSAVAAEIKQIRARRKTARMLMVVLLVFAICYLPISILNVLKRVFGMFA
                    HTEDRETVYAWFTFSHWLVYANSAANPIIYNFLSGKFREEFKAAFSCCCLGVHHRQED
                    RLTRGRTSTESRKSLTTQISNFDNISKLSEQVVLTSISTLPAANGAGPLQNW"
      exon
                    95,252..95,430
                    /gene="HCRTR2"
                    /number=3
      exon
                    101,753..101,996
                    /gene="HCRTR2"
                    /number=4
```

FIG. 1A

2/51

110,324..110,439 exon /gene="HCRTR2" /number=5 124,058..124,278 exon /gene="HCRTR2" /number=6 127,009..127,130 exon /gene="HCRTR2" /number=7 128,910..129,305 exon /gene="HCRTR2" /number=8

BASE COUNT 55,308 a 29,672 c 29,838 g 53,757 t

CGACTTGATTTTTTTTTTTGCATATGGATATCCAGTTTTCACAGCACTGCTTGTTACCCT CAGCAAAGAACAGTTGTCTGTAAATTCATGGGTTTATGTCTAGGCTCTCTGTTCTGTTCT ATTGGTCAACATATGGTCATATATCACTTAACTGCAGGGAAGGGATACATTCTGAGAAAT GCATTATTACATGATTTCATCATTGTGCAAACACTATAGAGTGTAGTTACAGAAACCTAG TATCTCTAGCTGTGTTCTTATGATTCAAATTTGCTTTGGTCATTTGAGATCCATACTGGT GGAGTCTAATTATTCAAAACTAGGGAAAACAGACAAACAGAAAAAAACTAAGACCAAGTTA CCCTTACCTTGACTAAAAAAAGAGACTCAAATAAATAAAATTGGAAATGAGACAGGAGAC ATTACAATTGATGTTAACAAAAAGATCATAAGGTACTATTATGAACAACTATACACCAAT AAATTGGACAACCTAGAAAAAAATGGATAAATTCCTAGAAATACACAGTCTATCAAACT GAAACAAGAAGAAATAGAAAGCCTGAACATACCAGTAACAACCAAGGAGACTGAGTAAAT AATCAAAAACCTCCCAAGAAGAAGAGTCTAGGACCAGAAGTCTTCACAAATGAATTCTAC GAATATTTTCAAACTCATTTTATGAGGCCAGCATTATTCTGATACCAAAACTACGCAAAA ATACTACAAGAACATAAAAACTACAAATGTGGGAATTATCATGTATACATATGCAAAAAT CCTCAGTAAAATCCTAGCAAACTAAATTCAACAGTACATTAAAAAGATCATATAGCATGA CCAGTGAAATTTCTCCTTAGGACGCAAGGATAAGTCAACATATAAAATTGAATGTGATAT ACCACTTTAACAAAATGAAGGATAAAAATCATATGATCATCTGAATAGATGCAGAAAAAG CATATAACAAACTTTGACGTTGTTGAGAAATTGAAAGCTTTTCTCTAAGATCAAGAACAA AGCAAGGATGCCCATTCTTGCTTCTATTCAGCATAGTGCTTGAAGTCCTAGTCTGGACAA ATAAAATCCACCAAATTGGAAAGGAAGAAGTGAAATTACCTCTGTTTGTAGATGAGCTGA ATAGCTAGAGCAAAGAATGAATTCAGTACAGTTGCAGAATGCAAAATCAGTATACAAAA AGTACTTGTAATTCTATATAATAGCAACAAACTATTTCATAAGGAAATTAAGGAAACAAT CCCCATTACAATAGCATCATAAATAATAAATCTTAAGAACAAATTTAACCAAGGAGGTGA AAGACTTGTGTACTGAAAACTATAAAATGCTGATAAAAAAATTAAAGAAGATACAATAAA TGGAAGATATTCCATGCCATGGTTTGGAAGAATTAATATTGCTAAATGTACATACTACCC **AAATGAAAACACAATCTTAACACTATTTAAACCAATTAAACAAACCTATGATTTCAATTT** GGTCAAATGTGTTAGAATGGATTTCCTTTTATTGTTTTTGAACTTGTCTCTTCCAAATTTC AAAGCCTGGTTCCTAATTTTTACTTGAAATACCAAATAACAAACCCACTTAATGAGCTCT GAGCCAGTTTTAGTAGCCAAACTTGTATTTAAATAGTGTGTTACATATTTGCACAAAAAG CCAACGGAGTCTAAATCAACACTAATTCACATCATTACTAGCAATCTAAAACATCAGATG ATAATTTTGCTGTTGTCTTTCAGGCAAGATATTCAACCATTGGTATTAAATGTTTTATAT GAATGTGCGGTGTTTTATTTCAGAAACACTTCTCTGAATTCCCAAGGCCTAAGAGCTATT CATCATAGAGGTTTGTGGAGGCGGTAGTTAGACATTTTCTACATGCATAATGTTAATTCA TTCAAACATTATAGAAAAAAGTTTGTAAAGAAGTTAATTTTCAAGGTGACAAAAAAATC AGATTGAATCATGTTTATTTTATTTCAATTTAAACTCGTTGGCTATCTTAGGAAATTCAC ATTGTTTTTGAAGAATATATGAACAAAGTTTGATTCATCTTATCTATATAAGCATGAGAG

FIG.1B

AAATACCTAGATGAGAGTGAAAAATGACTAATTTTGTGACCATTGTTATATCATAGATTA ACTTGTTCTCTACTTCTAAGCTGTGTGATCTTGAAAAGTCATCTAAACTTCAGGTAC CATCCTCACTTGCAAAATGAGGGGAAAAACCCCAGCACCTTTAATATGGTGTTATGTGGA TGAAATAAGTTAATATATTAAGTGCTTAGGTTTCATGCACTTTCTATATAGTATTAAT **AATATTATTGTTACATTATTATAGTTACATTATTATTTTTTATTAATATTATTGGAACATG** AATGGAATTGTTGTGGCTCATTTTAAAGATGCTGCAATGGAGACCAAGAGAAATTAAGTA TAATAATCCTAGTTAAGGTACAGCCATTTCTAATTACATTTTCCAACTGCTGCTTTTACT AGAACACATAATAATCTTATTTAATTATCACAACAATTTCTGTGGAGTTACTATTAATCC AGAGATGAAGAATCTAAAACTCTAAATTATCAAGCAACTATTCCAGCTTTAAACAACAGT AAAACTGGAATTAAAACTAGAGTTTCTTTATGAGGCCAGTATTACTCTTATACTAAAGCA CAAGCACACACACACACCAGATCAATATAACTTATGGATGTTGATGCAAAAATTTT CAAAAAATTAATAGCAAATCGAATCCAGCAGTATTTTAAAGGACTATACACCATGGACAA ATGGGGTTTATTCCTGGGATATAAAGTTGGCTAAACTTAATGAAAATCAACCAGTGCAAT AAATAATAGTAATTAAAAAAACATAATTATCTCACTAGATGTACAAAAAATGACAAGATC CAACATGTTTTCAATATAAAAGCATTCCACAGACTAGGAATAGAAGGGAACTTCCTCAAC TTTACAAAGAACATCTACAACGAAACCACAGCTAACATCATATTTAATGGTGGAAGACTG AAATCTCTGAATATTTTCCCCTAAGATCAGAAAAAGACAAAGAGGTCTACTAATTCTATT CAACATTGGAAAGATAGTTCTAGTCAATTAATCATTAAAAAAAGGCATTTAGATTAGAAA GGAAGTAAAATTACCTCTGGCAGATGACATAATCTTATACATGGAGAACTCCTAGAGATT ACATACACACTCACAACTACCAGAGTTAATAAATGGGTTCTACAAAGTTGCAGGATACAA TATCAATTCTCAAAAAACACTTGTATTTCTATACACTAGCAAATAACTCTGAAAATGAAA TTACCAAGAATGTACAAGTTTTATGTACTGAAAAATAAAAATGTCATTAAAAATAGTTA AAGAGAATCTAAGCACATTGCAGTTTTCTGACTCCAGGCCCGGGCTCTTGGATGGCATCT CTGGATCCACTCAGGACCAGGGAGAACTTGTTGCCCTGAAGGGAAGGACACAAGTCTGAC TGGCTTTACCACCTGCTGATTGTAGAAGCCTAGGGCCTTCAGGGAACACAGGTGGTAGCC AGATAGCAGTTACCATGGGCATTAGGCATGACCCAGTGCTATGTTGGCTTCTAGTTTGAC CCAGCACAGCCCAAGGGTGGTAACCACATGGGTGCTTGTGTCACCCCTCCTTAAGTTCCA GGCAGCCAGCAAAGAGAGAGTGACTCTGTTTGGGAGAAAGTAAGGGAAGAAGAATAAAAGT CTCTGTTGGTAATACAAGGAATTCTTCCAGATCTTATCCAAGACCTCTATGAATCTGCAA CAGCCAAAGCATTATTAGTTTTCAGGTTTCCCCAGTGCAGATATGACTGCAATGATCAAA GCACAAACAAGTTTGGACTGGGAGGACTACAATAAATACCTAACTTCTCAATGCCCAGAA AGGAACAGCTCTGGTCTGCAGCTCCCAGTGAGATCAGTGCAGAAGGTGGTGGTTTCTGCA CAGAGGGTGAGCCAAAGCTGGATGGGGTGTCACCTCACTGGGGAAGCACAAGGGGATGGG GAACTCCCTCCCCTAGCCAACGGAATCTGTGAGGGACTGCCATGAGGGATGGTGCATTCT GGTCCAGATACTATGCTTTTCCCATGTTCTTCACAACCCTCAGGCCAGGAGATTCCCTCG GGTGCCTACACCACCAGGGCCTTGGGTTTCAAGTACAAAACTGGGTGGATCTTTGGGCAG GCACCGAGCTAGCTGCAGGAGTTATTTTTCATACCCCAGTGGTGCCTGGAATGCCAGTGA GACAGAACCATTCACTCTCCTGGAAAGGGAGCTGAAGCCAGGGAACCCAGTGGTCTAGCT CGGTGGATCCCACTCCCATGGAGGCCAGTAAGCTAAGATCCACTGGCTTGAAATTCTCAC TGCCAGTGCAGCAGTCTGAAGTCAACCTGGGATGCTTGAGCTTGGTGGAGAGAGGGACGT CCACCATTACTGAGGTTTGAGTAAGCAGTTTTCCCCTCACAGTGTAAACAAAGCCACTGG GAAGTTAAAGTAGGTGGAGCCCACGACAGTTCGGCAAAGCCACTATAGCCAGAATGCCTC AGCTTATAGATCAAACTCCCATCTCCCTGGGGACAGGGCACCTGGGGAAAGGGGCAGCTGT GGGTGCAGCTTCAGCAGACTTAAATATTGCCGCAAGCTGACTCTGAAGACAGCAGGGGAT CTCCCAGCACAGCGCTCGAGCTCTGCTAAGGGGCAGACTGCCTCCAAGTGGGTCTCTG ACCCCTGTGTCTCCAGACTGGGAGACACCGCACAGCAAGGGTCGACAGCACCTCATACA GCAGGCAGCAATCTTTGCAGTACTGTAGCCTCTACTGGTGATACCCAGGCAAATAGGGTC TGACGTTGACCTCCAGCAAACTCCAGCAGACCTTCAGCAGACGGGCCTGAGTGTAAGAAG GAAAATTAACAAACAGAAAGGAATAGCATCAACATCAAAAAAACAAAAACAAAAACAAAA

FIG. 1C

ACAAAAACAAAACAGCACATCCGCACAAAAACCCCATCTGAAGGTCACCAACACCAAAT ACCAAAGGTAGATAAATCCACAAAGATGGGGAAAAACCAGCACAAAAAAGCTGAAAATTC CAAAAAACAGAATACCTCTTCTCCTCCAAAGGATCACAATTCCTCACCAGCAAGGGGACA AAACTGGACAGAGAATGAGTTTGATGAATTGACAGAAGTAGGCTTGAAAAGGTGGGTAAT AAACTCCTCTGAGCTAAAGGAGCATGTTCTAACCCAATGCAAGGAAGCTAAGAACCTTGA AAAATGGTTAGAGTAATTGCTAACTAGAATAACCAGTTTAGAGAAGAGCATAAATGACCT GATGGAGCTGAAAACTATAGCACAAGAACTTCGTGCAGCATACACAGGTATCAATATCCA AATCGATCAAGCAAAGAAAAGAATATCAGAGATTGAAGATCAACTTAATGAAATAAAGTG TGAAGACCAGATTAGAGAAAAAAGAATAAAAAGGAATGAACAAAGTCTCCAAGAAATATG GGAATATGTGAAAAGACTAAACCTACATTTGATTAGTGTACCTGAAAGTGACGGGGAGAA AGGAATCAAGTTGGAAAACATTCTTCAGGATATTATCCAGGAGAACATCCACAACCTAGC AAGACAGGCCAACATTTAAATTCAGGAAATACAGAGTACATCACAAAGATACTCCTCGAG AAAAACAACCCCAAGACACATAATTGTCAGATGCACCAAGGTTGAAATACAGGAAAAAAG TTAAGGGCAGCCAGAGAGAAAGGTCGGGTTACCCACAAAGGGAAGCCCATCAGACTAACA GTGGATCTCCCTGCAGAAACCCTACAAGCCAGAAGAGAGTGAAGGCCAATATTCAACATG CTTTAAGAAAAGAATTTCAACCCACAATTTCATATCCAGCCAAACTATGCTTCATAGTG AAGGAGAAATAAAATCCTTTACAGACAAGCAAATGCTGAGAAATTTTGTCACCACCAGGC CTGCCTTACAAGAGCTCCCGAAGGAAGCACTAAATATGAAAAGGAAAAACCAGTATCAGC CACTGCAAAAACATATGAAATTGTAAAGACCATCAACACTATGAAGAAACTGCATCAACT AATGGGCAAAATAACCAGCTAGCATTATAATGACAGGATCAAATTCACACATAACGATAT TAACCTTAAATGTAATAGGCTAACTGCCCCAATTAAGAGACACAGACTGGCAAATTGGAT AGAGAGTCAAGACCCAACAGTGTGCTGTATTCAGGAGTCCAATTCATGTGCAAAGATACA TATAGGCTCGAAATAAAGGGATGGAGGAATATTTACTAAGCAAATGGAAAGCAAAATAAA GCGGAGGTTGCAATCCTAGTCTCTGATAAAATAGACTTCAAACCAACAAAGATCAAAAGA CTATAGAGACTTAGACTCCCACGTAATAATAGTGGGAGACTTTAACACCCCACTGTCAAT ATTAAACAGATCAATGAGACAGAAAATTAACAAGTACATTCAGGACTTGAACTCAGCTCT ATTATTCTCAGCACCACATTGCACTTATTCTAAAATTGACCACATCATTGGAAGTAAAAG AATCAAATAAGAGCTCTGGAATAAGAAACTCACTCAAAAACCGCACAACTACATGGAAACT GTTACTTGAAACCAATGAGAACAAAGACACACATACCAGAATCTCTGGGACACAGCTAA AGTAGTGTTTGGAGGGAAATTCATAGCACTAAATGCCCCACACGAGAAAGTGGGAAAGATC ATAGATAGATCACTAGCCAGACTAATGAAGAAGAAAAAGAGAAGAATTGTATAGACACA ATAAAAAATGATAAAGGGGAGATCATCACTGATCCCACAGAAATACAAACTACCATCAGA GAATACTATAGACACCTCTATGCAAATAAACTAGAAAACCTAGAAGAAATGGATAAATTC CTGGACACATACACCTTCCCAAGACTAAACCAGGAAGAAGTCAAATCCCTGAACAGACCA ATAACAAGTCCTGAAATTGAGGCAGTAATTAATAGCGTTCCAATGAAAAAAAGCCCAGGA CCAGATGGATTCACAGCCAAATTCTACAAGAGGTACAAATCAGAGCTGGTACCATTCCTT CTGAAACTATTCCAAACAACAGAAAAGAAAGACTCCTCCCTAACTCATTTTATGAGGCT GGCATCATCCTGATACCAAAACCTGGCAGAGACATACACACAAAAAAGAAAATTTCAGGC TAATATATCCCTGATTAACACCGACGCAAAAATCCTCAATAAAATACTGGCAAACCAAAT CCAGCAGCACATCAAAAAGCTTATCCACCACGATCAAGTTGGCTTCATACCTGGCATGCA AGGCTTGTTCAACATACGAAAATCAATAAATGTAATTCATCACAAAAACAGAACCAATGA CAAAAACCACATGATTATCTCAATAGATGCAGAAAAGGCCTTCAACAAAATTTAACAGCC CTTCATGCTAAAAACTCTCAATAAGCTAGGTATCGATGCAATGTATTTTAAAACAATAAG AGCTATTTATGACAAACCCATACCCAATATCATACTGAATGGGCAAAAGCTGGAAGCATT CCCTTTAAAAACTGGCACAAGACAAGGATGCCCTCTCTCACCACTCCTATTCAACATAGT AAGAGAAAGTCAAATTGTCTCTGTTTGTGGATGACATCATTGTATATTTAGAAAACCC CATTGTCTCAGCCCAAAATCTCCTTAAGCTGATAAGCAACTTCAGCAAAGTCTCAGGATA CAAAATCAATGTGCAAAAATCACAAGCATTTCTATACACTAATAATAGACAAACAGAGAG

FIG. 1D

CCAAATCATGAGTGAACTCCCATTCAAAATACCTAGGAATACAACTTACAAGGGATGTGA ATGCAAAAACATTCCATCCTCATGGATAGGAAGAATCAATATCATGACAATGGCCATACT GCCCAAAATAATTTATAGACTCAATGCTATGTTCATCAAGCTACCACCGAATTTCTTCAC AGAATTAGTAAAAAACTGGCCAGGCTCAGTGGCTCACGCTTGTAATCCAAGCACTTTGGG AGGCCAAGGCAGGAGGATCAAGAGGTCAGGAGATTGAGACCATGGTGAAACCCCGTCTCT ACTAAAAATACAAAAAATTAGCCGGGCGTGGTGGCAGGCGCCTGTAGTCCCAGCTACTTG GAGAGGCTGAGGCAGGAGATGGCGTGAACCCAGGAGACGGAGCTTGCAATGAGCCAAGA ACAAACAACAAAAAAAAAAACTACCTTAAATTTCTTATGGAACTAAAAAAGAGCCCAT ATAGCCAAAACAATCCTAAGCAAAAAGAACATAGCTGGAGGCATCATGCTACCTAACTTC AAATTATGCTACAAGGCTACAGTAACCAAAACAGCATGGTATTGGTATGAAAACAGATAT ATAGACCAATGGAACAGAACAGAGGCCTCAGAAATAACCCCAGACATCTACAACTCTCTG ATTTTTGACAAACCTGACAAAAACAAGCAATGGGGAAAGGATTTCCTATTTAATAAATGT TGTTGCGAAAACTGGCTAGCCATATGCAGAAAACTGAAACGGGACTCCTCCCTTACACCT TATACAAAATTAACTCAAGATGGATTAAAGACTTAAACGTAAGACCTAAAAACCATAAG AACCCTAGAAGAAAACCTAGGAAATACCATTCAGGCCATAGGCATGGGCAAACACTTCAT GTCTAAAACATCAAAAGCAATGGCAAGAAAATCCCAAATTGACAAATGGGATCTAATTAA ACTAAAGAGCTTCTGCACAGCAAAAGAAACTATCATCAGAGTGAACAGGCAACCTATAAA ATGGGAGAAAATTTTTGCAATCTGTCCATCTGATAAAGGGCTAATATCCAGAATCTACAA TGAACTCCAACAAATTTACAAGAAAAAAAAACAACCCCATCAAAAAGTGGGTGAAGGATGTG TCATCACTGGTCATTGGAGAAATGCAAATAAAAACCACAGTGAGATACCATCTCACTCCA GTTAGAATGGCGATCATTAAAAAGTCAGGAAACAACAGATGCTGGAGAGGATGTGGAGAA ATAGGAACGCTTTTACACTGTTGGTGGGAGTGTAAATTAGTTCAACCATTGTGGAAGACA GTGTGGTGATTCCTCAAGGATCTAGAACCAGAAATACCATTTGACCTAGCAATCCCATTA CTGGGCATATACCCAAAGGATTATAAATCATTCTATGATAAACACACATGCACATGTATG ATAGACTAGATTAAGAAAATGTGGCACATATACACCATGAAATACTATGCAGCCATAAAA AAGGATGAGTTCATGTCCTTTGCAGTGACATGAATGAAGCTGGAAACCATCATTCTCAGC ACAATGAGAACACATGGACACAGGGGGGGGGAACATCACACACCAGGGCCTGTCAGGCAGT GGGGGCTAGGGGAGGGATAACATTAGGAGAAATACATAATGTAGGTGACAGGTTGATGG GTGCAGCAAACCACCGTGGCACATGTATACCTATGTAACAAACCTGCACGTTCTGCACAT GTATCCCAGAACTTAAAGTATTAAAAAAAAAAAGACCATTTATGAAAACATGACCTTACCA AAGAACTATATAAGTCACTGGAGACCAATCCTGGAGTGACAGAAATATGTGACCTCTCAG ATGGAGAATTCAAAATAGCTGTTGTGAGGAAATTCAACAAAATTCAAGATGACATGGCAA AGGAATTCAGACTTCTATCAGATAAATTCAAAAAAGAAGATGAAATAATTTTTTTAAAAA TTCATGCAGAAATTTTGGAGCTGAAAAATTCAATTGATATACAAAAGAATGCATCTTACC AGCAGAATTGATCCTGCAGAAGAAAGAATTAGTAAATTTGAAAACACTCTATTTGAAAAT ATACAGTCAGAGGAGACAAAAGAAGAAAATTAAAAACAATGAAGCATACCTACAGGATCT AGAGAGAGAGTGGGATAGGGGTAGAAAGTTTATTCAAAGGGATAACAATAGAGTATCAGT ATTCAAATACAAGGTTATGGAACACCATTCAGATTTAACCCAAAGAAGACTACCTCAAGA CATTTAATAACTGAACTCTCATTCAATGGGAAAAGTAAAGTCCTTTCAATAAAGGTGTTG GGATAATTGGGTATGCAAAAAATGAATTTGGATACCTTTCTTGTGTCATATACATAAAAC CACAGTAAATTTTTGTGACCTTTGATTAGGCAATGATTTCTTAAATATGATAAAATATGG TAAAAGCAACAAAAGAAAACATGAATAAATTGGATCTTATCAAAATTTAAAACTTTTTTG CATCGTAGAATACTATCAAGAGTATGAAAAGAAAACCTACAAAATAGGAGAACATGTTTG TATATATATATATATATATATACTCTTACACCTCAACTATAAAGAGACGAATAACCCAAT AAGTTCATCAAAAGATGCTCATCATCTTTACTCAGGAGGCAAATACAGATTAATATTACA ATGATATTAGACATGGATTTGTCATATACAGACTTTATTAAGTTAGATTCCCTCTATGCC TAATTTGTTGAGAGTTTTTATCATGAAGAGATGTTGCATTTTGTCAAATGCCTTTTCTGT GTCTTTTGAGATGATCATATGGTTTTCGTCCTTTATTTTGCTGATATGATGTACCACATT

FIG. 1E

TATTGATTTGCATTTATTGAATCATCCTTCCACCCCTGGGATAAATCCCACTTGATCATG GTGTATTATCTTTTGATGTTTTTTGGATTCACTTTGCTGATATTTTGTTGAGGATTTCT GCATCTATAATCATTAAGGATATTGGCCTGTAGTTTTCTGTTTTTATGTTGTATTCTAGT CTGATTTTGGTATCAGGGTAATGCTGTTCTTGTTGAGCGTGTCAGGAAGTCCAAAAGACT TAGAATTCAGCAGTAAAGCCATCCAGTTCTGGGCTTTTCTTTGTTAAGAGACTTAAAACA CACACAACGCACACAAAATGAAATATCACTTTCCACCCATTATAATTTACAAAGTGGA AAATAACTCGTGTTGATAAGAATGTGGAAACCTTGAAACCTTCATGCATTGCCAGTGGTA ATGTGAAAGAATCTTGCCATTGTGGAAAACAATTTGTCAGTTCCTCAAACAGTTCAACAT AGAGTTACTGTATGAAATAATTCAACTCCCAGGCATGCACCCAAGAGCATTGAAAACATA AGTACACAAAAACTTGTACAAGAACAGTCAGATCAGTATTATATAAATTGGCAAAAA ATGGAAACAATCCAAATATTCATCAACTGCTGAATAGATAAAATGTGGCATATCCATATA ATTAAATACTATTCAGCCACAAAAATAATAAAGTACGGATAGACACTAAAACATGGAAGA ACCTTGAAAATATTAAGCTAAGTGAAAGACATAAGGACACAAAAACCCAACATTTAAAGGAA ATTTCCAGAATTGTCAGATCCACTGAAGAAGAAACTTGAGTGTTTGCCAGCATGTGGGAG GAGAGGAAAATCAGTAGTTATGAGGTTTCTGGAATTAGTAGTGCTGATGGTGACACAACA TTGTGAATATACTATAAACCACTAAATGATACCTCTCAAAATGGTTAAAACATTACTGTT GTGTTATGTGAATTTACCTCAATTAGAAAAGAAAAAATCTTATCAATAACAAAGAGAA ATTTCCACACAAGGTGGGATCGCTTCCACAGTGCTACTCAATGCAGTTTAGCGATTGCAT TTGTATTGGAGTAAAAGCATGTCACATTGCTTTTAACATTGGAGTCCAATACATAAACCT CTTTCACCATAACTATATGGAGTTCATTGTATGTATATTTATAAAATGGAATTAAGATG AATTTCACAACACAATGGATCATTTTTTTTTTCATGTGGAAAATCAGAACACATGCCTTA ATGGTTACATGCCCCACCTGCTGCTCACCTAAAAGTAAATTTCCTCTAACTCAGACAAAT ATGTTATTTTCAAGGAAAAGAAGCCCAGAGAACTGAGATCCAGAAGAAATAACATGTATT GAAAGCACACAGAAGTATTTCAATGAACTCAAACCCAAGATTGTAGAAAACTCTCATGTG CCCACCCTTCAGAGCACCCGACGATAATGGATAGTTTCTAGCAGGGTGTCTGGAATGGGC AAGTACCCCCAAAGTTATAGTTTGTACTGCAAGACTTGAACCCACTCTTTTTCTGCCCTC ${\tt TATTATTATTTTGCATTTTAACCATTTATTATTTTGAAAAGAAAAGAGAATTTTTAGAA}$ TATGGAAAGAGGAAGTGAATTAATAAAATAGCACACCCTACATAGAGACTGCTAATCCAT CTCCAGTCTAAAGATTTAGTAATAGGCAAGAATATACATATCCAGGAATTTCCTTGGTGT TACATAAACAAAGGCGGCACATATGTATATTTTCACAAAATATTCACTGTTTGAAGAAG GAATTACTCCCTTCAATTGAGTTCAGGCCTGATCAACAAGTAGTGATTGGCCAACAGCTA GGAGATGTCATCCTGAAGAGTATAACAAGTTCCCCTATAATTCTACTTTTCAGTACTGTT TAAAATACAACTGGATTTTTTTAAATATGTAAAATTTATATAATTTTACAAATGTCTTTG TTAAGAATTAAAACTATCATTAGTAAAGGACACAGCTGGAAAATTGAAAACATTTTGGTT CTCTACTGTGGAAACAGAATAGAGTAACAGCAAAAAGCGTATTTCTGGAATTGGACCCTG ACAACTCTGCTTAAACACTCCACCACTTTCTAGCTATATGACCTTGGGTAAGTTACTTAA CTTCTTTGTGTGTCAGTTTCTTCATTTGTAAAATTGGAATAATAGATGCTTTTTTTGAGA ${\tt CAGTGTCTCATTCTGTTGCCCAGGCTGGAGTGCAGTGGCGTGACCACAGCTCACTGCAGC}$ CTCAACCTCCTGAGTTCAAGTGATTCTCCAACTTGAGCCTCCCAGATAGCTAGGACCACA GACACATGCCACCATGCCTGGGTAATTTTTTTTTTAAGTTTTTCATAGAAATAGTGTCTC ACTAAGTTGCCCAACCTGGAAAATTGGAATAATAATTCATAAAATCTTCCTCCTAGATTT GTGAAGATCAATTGAGTTAATGTAACGTACTTGGCACAGAGCTTGGCCCATGTAAT CAATGTCTTTTCCATATGGTTTCATTGACGCCACTTTGGGAAAATAGATGTCTCTTCTGC TTGCATTTTCAGACCTTTTTAGGTGTATACCTTAGGGCATTTGCTTTACTGACCAAAATT ATTTGCCGGCTACTCTGTGCTTTTCATGACACACTGAATAAGACAGGAAGAGTGTTTATC TATGCTCAACATAAGATAGGCATATAATGGAAGCTTCGTATATATTTGTTGAATAAAAA CATAAGGGGAAAATATCAGATCTAATAATGCAGGACAGGAGGCAAGATGGAACGGAGAGA ACCTTGTCTGAGAAGAGACATAATTAAAACAGGGCATGGGAGGTAATAGAAAGATTGGAG GAAAAAGAGACAGAGAGACAGAAATGTTTGTGGTAATTTGTGACAAGTAGCTTTGATTGT TCATGGCCTAATCTTTTAGGGCATGAGGTTATTTCATTCTCTGTAGCCCACCGAGAGTGC GTACAGTGACACATGTTATGTAAGTCCCCTTTTCCCTTTTTATAAATGTCTAGACCCCCT

FIG. 1F

GTGATTTGAGACTTTTCTAGAAGAATTTAGCTGAAGACCATATTGTTTTTTAAATGTAGT ATATAAGCTCAGTATCATCATTACCAACAGTGCTCAGACTTGATTTTATTTTCATTCCAA CAGCAAAGGAAAGCAACTTCTTTCATGCTTCCATGCCACTCTGCATCTCTCTACCT TCACAGAGTTTCTCAATAATGGCAACATTTCCAGTTCACCAATGGACTGAGAGATCATTG AGGCTAGACTAGTCTTATTAATCCTTATACCCCAGCTCCTAGCCGAACTCCTGGACACAC AATAGATACTCAGATACATTTACTGAAATGCATATAGAAAGTTACACCTGCAAAAAAGAT GATCTCTCACCAGGAATAAGAAAATATAATCTGGGACAGCCCATATATGAGATCTCTAAA CTAGGTATTTAAACAGAATTATTCTGAATGTTGTGAGCTACATTTCTTTTTTACCTTTTA CATAGTATTTGTATATTTTATAGGGTACATGTAATATTTTGTTACACGCATAGAATGTG GCTAGGAGCATGTTAAGTCCTCTTTTTAGCTATTTTGAAATGTACATTGATGTTAACTA TCATTAACACAGAGTAATTGATATGTATAGCAAATAATATTTGCAGTAGGATATCACATG TTTACTTATTTATTTATTTATTTTTTTTTATTATACTTTAAGTTCTAGGGTACATGTGCA CAACGTGCAGGTTTGTTACATATGTATGCATGCGCCATGTTGGTGTGCTGCACCCATTAA ACAGGTTCCAGTGTGATGTTCCCCTTCCTGTGTCCAGGTGTTCTCATTGTTTAATTCC CACCTATGAGTGAGAACATACGGTGTTTGGTTTTTTTGTCCTTGCGATAGTTTGCTGAGAA TCATGGTTTCCAGCTTCATCCATGTCTCTGCAAAGGACATGAACTCATCCTTTTTTTGGC TGCATAGTATTCCATGGTGTATATGTGCCATATTTTCTTAATCCAGTCTATCATTGTTGG ACATTTGGGTTGGTTCCAAGTCTTTGCTATTGTGAATAGTGCCGCAATAAACATATGTGT GCATGTGTCTTTATAGCAGCATGATTTATAATCCTTTGTGTATATACCCAGTAATGGGAT GGCTGGGTCAAATGGTATTTCTAGTTCTAGATCCTTGAGGAATTGCCACACTCTCTTCCA CAATGATTGAACTAGTTTACACTCCCACCAACAGTGCAAAAGTGTTCCTATTTCTCCACA TCCTCTCCAGCACCTGTTGTTTCCTGACTTTTTAATGATCGCCATTCTAACTGGAGTGAG GCACTGGTCTGAAAATATCAATTCATTTAATTCTTTTAACAACCTTAAGGGGATATCATG CAGCAGAAATATTTGAATTGAAGAGAAGAGTAATACCTAAGAACTAGAAATTCCTTTCTT ATGTTTCAAAAGATATCAAAAGATCTAAGGAAGATATTCACATCAAAAATGAGTATTATA ATATTTATCTATGGTGCACTTGCAAAAAAGAAAACAAGTAATAATCTGAAGATTTAA GTGAATATTTTATGACATTGGAGTACCACATATTTAGAAGAAAGCACCAGAGAAATCATA GATAGAAGGAAATGGAATATTTGTAGGATCAAGATAAATACAGCTTGTCATAAAATAAAG ACTGTGATGATTAATTGTAGGTGGAAGATTTACGAAGAGAGACTGAAGTATAGACAAGT TGAAGTGCCACAAAATGAAAGCTAATGACACTGACTACTTAGGAAATAGCAGACTGGGTC CATATTTATAGATTGTCAATGACAAGGAATTTGCAGATGTTAATGAATATAGATCCGAAC TTAAGTTGCAACAACCTTTCCCACTTTGAGATGAATAGTGCATGGAAGAGTAAAATGCAG **ATGTTAATAAATCAGAGGAAGACATCGTGCCAGAGTATAAAGTTGACAGATTTATGCCGA** TGAACTTGAACAAAGCCACAGAAGGCCTACTTGTCAAATTTACTGGTGACAACAGGTCTG GAGAAATGGCTAATGTTTTGGATAATAGCATTAGAATTTAAGGTCTGTTTAAACTTCAAA TTAACAGAATGAAATTAATATATGCACATATCAATTGGGTCTTTTGCTTATATATCATCT CTTAATAGAGCCTTTTTGAACAATCATTTCTAATGTGACCTTTGGGATTTTCTACTCATC ATCACCTCATCCTGTTTGGTTTGCATTATAGCATCTATCCCTTCCTAACGTTTTCCCTAT GTATTTGTTAGTTTGTTTTTTTTTAATCTAACTTTACTAGAAAGTAAAATGCATGGAAAC AGCAACCTGTTTAACTTTGTATCACTAAGAGTGGAAAAATAACCCTCAGGAAATATTTGG TAAAATAATAAAATGCCCATTGATGCCCTTCTCTTAAAAAGAAATTTAATTAGTGCAGAT TGGGGAAATACAACAATATTTCTCATAAAATGTGATATCTATACAATAACAGAAGTACTA TGTCCCAAAAGTATTCTATAAATAGAAGAAGAACAGATGGTTTTGCTGCTGATTAATC CATTTATCTTTCGTAAATCATCTAATTTCCCCAGGAACAGCTTCCTCATCTATTAAAGGG GGTTAGTAATAGCTAAGCCCTCAGGGGTTTAAAAATGCATATGAAATAATTTTATAAACC ATAAAGCACAAAACAAATATGAAAAATTATGATTGGAGGAGGGGGTGGGGTAGTTAACTA AATCTCAGTGTAAACCACCAATGTCTTGTGTGTGTTGAAAAAATAATTACATATAAAAAC TGGTTGCATCCAAAGAATAATGTACTTTTTGCACTGGCAAGACTCAAACCATATTATTGT

FIG. 1G

TACTTCCTCCCAGTTACATATTTTGCAAGATATTGACAATTGTCTAAAGGAAGACCAAAC AGATGTAGGTGGGAGCTACTGTCATTTGAACAACATTGAAAAGAAAAATACTAAAAAAGA AACATGAGGGCATATAAAGGAGCGCTGGGGCTGTGATGTTTATTTTGAATCTGTGAAGCA TTGTCATGTGGAAGATTTATTCTGTGTAGCACCAAGATGCAAACTAGGAATTAGAGGTAA AAGTCTCAAAAAGACAAATCGTGGCTTGAGACCTTGGTTTAATGTAAGAAACAGTTTTCT CACCCTTAGAGCACTCCCATAAGGATGGAAGTAGTGAATTGTGGTGGTCACATTCAAGCT AGATGGGGACATGTCAGCAATGTTATCAGGAGGCTTCTACTCTGAAGCTGAAGTTCAGAC AAGATTTCCAGGCTCTTCCCAAGTGCAAGATTGTAATTACTTAAATGCAATATTTTTACC ATGTTTATTAAGAATAAAAGGATCATGAATTCACATTCTGACAAATGCTAGAATACTTAT TATTAGAGACAAAACCAGTGCATGAGAGAATGGCAGGTGACATCAGCCCTGAATCAATGG CATCCTACAGTGAATGTTTAATATCATTGAGTATATTGGTGGTCTGTCATGCTTGACAAC ATTAACTATGATCATATTTATGACACTTGGCGTCCTTCAAGAATTTGTAGCTCTATTTCA CATGACACTTAACTATCGCAAATACAAATTCCAGCTAAATAGACCCTTCAGTTTAAAAAC AGTCTCATTCTCAAATTTTAAGGAGAAAGTGAAGACGGAGATGTCTTAAAGACTCGGCAA GTACTAAGTTGGCAAATGTCAAATGTTAAAATAAGTTTATATTAAATGTTAAAGTGTTTG CCTGGAATGACTTTTCCATTGTCCTGCTTGAGAAACACAGAGGCACCTCCTTATTGCTTT TATATTTGCTTTACAAAGACAAATGTATCAACATGCTCTGTATTAATTGTATGTTGACAT TTTTGTCATATCCACAGACTGATGCATGTCTGTGCATGGTTTATAATAAGTGCACGTAAA AATAGAGAAAATAAGTAGAAAAAGAGAGAGATTTAACTCTCACCCCCACCCCCAAAAA CGAGGGAGGAGGCTGTGGGCTGCGGACTGAGTGCTGGAATGAGGAGTAATTGAGCTTCAG $\tt CTGAGCCGGACGTAGCTTTCTCCTCCTGGTGTCATTGCTGCAGCCTCCAGTGCCGGGTCC$ CTAGTTCCTCAGCTGCCTATCTTCCCGGTGCAACATCGCCTGTAAAGACAGCAAAGCCAC CGCAGAAGTTGCCCGGCAGAAGACTCCGGAGGCATTGGCTCAGTAACTTTTCACGTCATT TTCTGCTCGGGAGCCCCTTCTAGCCTCTCCGCGCAGCCTTTCCCACCGCAAATCACCAGT GCTCATGGGGCAGGCGGAGAGGAGCTTGCAGCATTGAGCGGAACCGGACTTGAGCCCGTG ATGTCCGGCACCAAATTGGAGGACTCCCCCCTTGTCGCAACTGGTCATCTGCTTCGGAG CTGAATGAAACTCAAGAGCCCTTTTTAAACCCCACCGACTATGACGACGAGGAATTCCTG CGGTACCTGTGGAGGGAATACCTGCACCCGAAAGAATATGAGTGGGTCCTGATCGCCGGG TACATCATCGTGTTCGTCGTGGCTCTCATTGGGAACGTCCTGGGTGAGTCTCCTCCCGGG CAGCCCTCCTAGGGGCTATCACCCCCTCTCCGCCCCGGGCTGAGAAGGCTCTAAAGAGAC CCCTCCCTCCCCGGGAAGCAAACAAAGAGGTCGCTGCTCTCGGATGGGGTTTTCTAATA AAATAATAATAATAGAAAGTTTTCTGATTTTCCGAACCGGGACCGAGCCCTGGAAAG GTTATTCCCTGTTTTGCAGGAATAACGGGGAAACCGCGTTTCTTTTTCGAGCACCTAGAT TACAAGCGCAGGGAGAGGGCCGCGGCAGGGATCTCCAGGTGGATTTTGTTGAGTGTGTG TGTGTGTGGGTGGGTGGGGGGGGTCAGTCATCCCTTTGTGTAACGTGGCTGGGTGTT TCAGGGGGGTTGGGACGAGACAGAGCTTGCAGAATACAAAGCTACATCCCTAAGGAGCAA GCTCTCTGTGGCTGTGGAAGTCACAAAGCATTTGTGAGCTAGGTGGCATTGCCCTTTGGC GAGGAGGTTTAGTCTCCAGTCAAGAGGTGGTAATGAACCAGCAGGAGTGGAGACGGAGG CAAAGCAGGGAAGTGCACTCACTCATAGAAGCTGAATTAAACAGGATCCATGCCTGGAGC ATTCTTACATCCATTCAGCCAAATATTTTTTTTTTTTCAGTCTGCTTGTTGCCAGGCTCAG GACTAAGCTTAATGCTAGGCTATTTGTCCCGGTCTAGGTCTGTATGCAAACACGGGTTTC CTCGACCCCTCATCCCCCTCCCCCTAAACAATTTCTGAGGGTTGGGGAGGGGGTGAGATG GCAACATGGTGAGTGCGATGATGGAATGTATTAGGGCAGTTGGGGAATATACCTCCAGAA AAGGGGCTTTGGAAGGGAGGGATAACTTGAAATAAATTGTGAATGGAAGGAGAGTGTACC TTGATGAATGAAGAGTAGAAGGCTGGGAGACTTTTCACATGCAGAGGGCAGTGTGGAGGA AGTCTCTGCTGAAAATGACAGGAGATGGAGGAGGCTAGGAGTTGCTCTTGATTTCATTT ATAAAAGAAGAAGAAGGTGAGTGAGGTGAGATAGGCTGGGAGGCTTTGCAGTCAAAAGCA AAGAACTTGTAGCTGCAATGGGGACTGACAAGGAAATTATCAGGCTTTCAGACTAACCTG ATTTTTGCCTTCTCCCAAGTGTGTTGGTCTGGGTAGAAATCATCCCGAGTAGTCTCTC ACCAACTCAGCAGGCAGAATAGATGATAGTATGTGAATGACAGGAGTTCTCCAGAGTGTT GGTAGAATGTTATTTGAGGAGACAAGAAACCTCTGAGAACTTTAGTACATTTTTAAATAT TATTTTTAGACTGTTTTCCTTTGGTTGATTTAAAAGTAAAAATAAAGGAAATCTTTTTGG GATACTAACAAAATGAAACAAAAGTGGAAATACACAAGATTAGGATTCTTGTTATAAGCA

FIG. 1H

TAATTCTGTTGATAATAATCCTAATCTTGCTTTCCTTCTTGTTACCCATCCTTAGGA TTACATCTCTTAAGACACATGGCTACCAGCATAGCAACATTTTACTGCATTATGCCAACA CTTATTGATAAGTGAATAATCAAAATTGAACATATATTGAGTACCTACTGTGTGCCAGAG CCCTTCATGTACATTCTCTCCCTTAAATATCAAAATAACCCACATTAGCCAGAAGAAGAA ACAAGACTTAGAGAAATAAATGACGTATTAAGGGACATAATTTAAATTCAGTTCCATTT TTTCTGACCTCAGATCCAGAATTCTCCATTGTTATTCCACTCTAGAGCTAAAAAGCATAT AGAGAATAGATTCTCTGCTCCTGATTGTCTGCAAGTTTATTAGATGTGTTCCTGTTCTCC TCTGCATCAACGCCCACTGCCAATAAAGTACAATGAGGGATTAATGGCACTGTCATTCTC TATTTGCCTTCCTTATTTTAATTTTCGGCTGAATCTTTGTGGTAAAATGTGCTCTTCTTT GTTGTTATTGCATTTTACCTTGCATAGACCTTGTAGTGAATAGTCTCCATATCCTAATT GCATAGTTTAGGGATACATGTTTGCTAGCCTGGGGAGTTTTAGTTTCAAGAAGGAAACAC $\tt CTCTACAGTAAGGCTACTTGTTTCATAATGTCAAGGAAGATAGCACTGTCCACAGCCCCA$ CTTAAAGAAACTGACAGCTATTTTCCTCAGGACTGAATAACACTGAAATCCTCTCTGGTT GAACTGAAATGCATTCTTTTCTGACATACTGCCTGAAAGTTGATGAGGTTTAGGTTTGAC ATTTAAACAAACGAGTAGTGTCGTTACTCACAGACAACTTCCTGCTCTTTGATGTCACTG TCAAATTTGCAAAATGAATTAGATTGAGAATTGCTTCTTTGCCCCTCTGGTATAAGTAAT TTTGCACATAGAGTGGTAGGACAGGATGTCACATGATTTATGCAAAATAAAGATGCAATA TTAAGTATGAAGGTAAAATACCACAGTGTAGGCAGCAGATGTAATCACTGAGCCTTCAGG TCCAGTCACCATTTGTACTTTCATATAACTGCTTGGAAAATCTCAACCTTTTTGGGCTTA CAAATATAATGCCATCAGTTAGAAGTCATCTTCTCCACAATGTCCTTTCATGAAGTGATG TAATAGGATATGCTGTGGGTAGCATAACAAAGTCTTGATTGTCCTCATCTCTTTTTCTTC TCCCCATAGTCCCTCTTTATCACTATGCCACCTCTCCACTCTCATATACTCCTCCCAAAG ATGGAAAGCAGTTTCCTGGGGGAGTAAAGTTTTAAATAGAATGTTATGAGTATTTACATT GTAAGGTTCTTTTTGGAAGGAGTATCTTTTCAGTATCTTCAGAATAATGCCACCTATAAC CTATTCCTAACTATGTCTTCTACTACAGCTAAGTAGATGTATCAACTTATTCAATTGGTA TATTGTGAGCATTATCATTTTTTTAAATTAGTGTGTATATCAGGGGAGCCTCTGGGGAAA TGTAAAGAAATGTGACTGATGTTAATTTTTACTCCTGATTCCTTGAATGACAATTGTAGG GAGAAATGTGTTCTAGTCAGTTTAAACATTAAGTACCTAGGGAAAATGATCAATTTTCTG CTTCTCATATCTGCATTCAAAGATATCATATGTTTCATCTGGTATGCTTCTGTCATATCT GTTGTTGTCTCCATATGGAAAATAGGAAAACATCAGTCTAGCTATGCTTCTTTGCTTCTTG TGTGCCATTAGCAAGTTATTGAACTATCCAAGTCAATTTTTTTATAATTACAAATTAAAG ATCGATAATGACTGCATTATAGAAATAGTATCAGGATATAATGTACGTATACCCTCTATA AAGACATATAAAGGGACACAGGCATATACATATTTTTCTTGACACATAGACACTAATTAA TGTCAATTTTTATCCCTTAATTTTCATGACTGAACTTTTTGTGATGTGGTGTATAGCCAG CTTCTGCCTTCATGGGCCAGTCTGTATCTCTGTAGCTCTTTATGGCCTCTGCCCCAGCCT TTTCCTTAATTGCATATTTTCCTAAAAGGTGTGAATAAAATGGTGTTGGCACACATTACT CTCCTTTTCCACACTAGCTCCACCCACCCATCTCCTTCATACTGATTGCTTAACATTGCC TTAAATTTGCCTGTGCTGGGTCCCATTCATTTAGAGTTTTTTGAGTTGTTAATAGGTTGTT TAAGTAATTTACCTAATAACAGTTTACCCAAGTTAGGTGTGTGGAATGGGGAAATATTTG TAATAAGTTTGCTTCCTACAGAGTTAGTCTTGTGTCAGATATGTAAGTGGTAGAATTGCA AGTTCATGTTACTCCTAAGCCTAGAGACATTTATTTTCTGCTTCTCCGAATGCCCATTTT AGTTTCATGGGTGTTTGTAAACCCATCCTTACCTACACAGGAAGCAAAAAGGGGTTATTT CTAAACCCTTTTTAGATATAGAAATAATACATCACTCATCTCGGCCAAGACTCAATAGAA TCATGAATAGTGACTGTAAAAGGTAATATTAACTATTAGGCTTTAAACCTATTGTGCATT TTAGTTTTAAAATGCAAACATGCTAATCTGAATAAGAATTAATCTGATGCCTCTACATTT ATCCCAATGATAACTTTTAAGATGGCTATTTCATAGATAACAGCAACATTTATCATGGAC AGACAATAATGAGAATAACATGTGCAACTGATAATTTAAATGCAATGAGTTATTTCTGTA TTTGAAAAATATATTTGGGAAATGGGATAATTAAAAAATACCAGTTTTCAAGAGACCAA ATCTAAAACTCAAACATAAACACAATGCTCCAGTTTTTAGAAAACTGTCTTGATTGTAGT AGTGCCTACATACTAAATTGTATCATATGATTTATATTAATTTTCCTTATTTTGTATTTT

FIG. 1I

10/51

AGATTATATTTGAAAATTTTCATGTACTGCAGCTATGTTAGCATCTCAAAGTCTCCATAT TCTCACTCCGCTCCGAAACATCCACTGCTGATGTTATTTAACTAGTGAAAGAAGATCCTT CCATGTTTCTTTATAGCATTCTGACATCTTCTCCACCCTAAGGAATGCTGGCTTTATT **AAGTATGTTTCAGTCAATGACATGTGATTGGTGAAGCTGACGGTATTTGTCTTCAGTTCC** TTTTTTCCCTGCAAAGGAAATTTGTTGAATATTTATTGGGTACTATATGCCAGGTACTAT ATGTCAGGCTCCACTTACATATACTCTATTGATGCCTTACAACAACTTATAATGAGAAG ATTAATAGGTTTTACAAATAAGAAAAATGAATTCAAAGAGCAATGCTAACTTACTCAAAA GTTTAGTCAGGCAGTAAATAGCAGCACTAGGTTTCAAATATGGATTTAACAAATTCCATG GTCCATGCTTATTCCATTACTTCATCCTGCCTCTTTCCTTAGCTTCTAACCCTGACTGGA GATGCATAGGCAAAAAGAGGAAGGAAGAGATACTTAGATGTGCCCTCTAGACAATTTACA GAGTTGTTTGGGCATGTTGCCATGCTGTTTTTCTGATAGACTACAGTTCTTCAGCTCTGA GGATGAGCTCATTTGATAAGCCAATCAAGGTCGGGCTAGGGTTACTTTACAAGAGAAAAT TTCAAGGTAAAATAGGTGCTGCCAAAAATGCTTTTACCTGTTCAGGGGGTTGACTCACTG GAAAAAAATGTTAGATAATTGTGGCCAAGGATTATTTTGTTATTGAAAGTGCTATTTTT AGACACAATTTGAGCCTGAGAGCCTAAACACTTAACACTTCACATAATCTACAGATATTT CTTGCTCTGTCACCCAGACTGGGGTGCAGTGGCACAATCTCGTCTCACTGCTGCCTCCAC CTCCTGGCTTCAAGCTATTTTCCTGCCTCAGCCTCCCCAGTAGCTGGGATTACAGGCACA ATCTTGGCTAGGCTGGTCTTGAACTCTTGCCCTTGTTATCTACCCACCTCAGCCTCCCAA AATGCTGGGATTGCAGGCATGAGCCACTGTGCCTGACGTGAACAGGTCAATTTCTATATC ACATCAGACATGAAATGACCTTTAGATACTGACTTTGAAAGAGTTTGAGATGCTATTGGA TGAAACACATGACCCATATGACCAGTCTTTTGAATTGCTGACTCTGAGTATAAAATGTTT TCATTTCACCTTTGTTCACAATGAGAAGTGATCTCTTAACCAAGTAAATGAATTAAATCG ATATTTAAAATAACATTAAATTTCTTGCCAGAAAAACTGTTCTTTCATAAACAAAAAACA AATTGCTCAAAATAAATGACTATATCTTTATTTCTAAAAAATGTTTAGAGATTATTATTA TTGGGTCTTTACAAGTAATTTGCCTTCAATACTAAACACATGAGAACAATGTTTAATATT TATATAGTATTTACTCTTCAGAAGATATTTGTCCATATTCTCTCAGTTATTCTTCAC AACAACATTATGAGGTAGGTCTTTTTTAATGAAAAAAACTCAAGTGCTTGAAGTGATTT AAAATCACTGTGGAAGAAAAGCATGGGCATACAGAAAAGCCAAGTGGTTGTGTCAGCT TGGGAAAAGCTTGCAAATTTCCTGTATTTCAAGAGGCCAGGATGAGGTGTAATTATCT TTTACTGGTCTTCAGCTATCCTGTCTTTGATATGTGATTGTGTCAAAACTATGAGGAAAA ACTCACATTAACAAACTTCATAAACTTGTTAAACATAAAATAATAATTTCGATGTTTTAA TTTACAGTAAGAGTTTATTCTTACAAGTCCTTAAATACCCAAAGTTCTTTCAGTTATCAT AGTCTTTTTCAGTAGACAGAAATCCATGTGGACTGTTATTGTTCTGAATAGCTAGGCTAT GCCATAGTAGCAAACAACCCTGAATTTTCATTGGCTTAGTATCACGAAAGTTTATTTCT TGCTCATTTAACATCTGAGGTGGGTTGGAGAGTCTCCTTCATCCAATGACTCACAGTTCA GGCAGCCTCCACATTTTGTGCACTATCCCTAAAAGGTGGACTCTGTGGTAATCAGTTTCC AATATGGCTTCCAATGACCGCCCCGGGCCCCGGCCCCACTTCCTGATAGTCACATCATC GTGTAGTCCCTTTGCATATTATGCCAGAATTGGTCTGGGTGACCAACAGCTCATAGCAGC AGTGAAACGATGTCACTTTCAAGATTACATAACAGGAGCTTACAGCTTCTGGCTCAAGTA CCCACTTTCTCTAGCTCTTGGATCTCTTCTTCTGGAGGAAGTAAGCTGCCTTGTGGTG TAACAACCTCATGTGTGAGCTTGGAAGCAAATCCTTCAGACCAGGTTGAGTCTTGAGGTG ACTACAACAGCCACTACCCCAACCCACCCCAGCTTCAGTGCAACTTAGTAACAGACACT GAGTCAGAACTATTCAGCTAAGCTTCTTGCAGATTCCTGACCATTCAGAAGCTATGTCAT AATAAATTTTTGTTGTTTGACTTCAGTTTCGGGATAAGTTGTTGCACAGCCTCTAAAGTT GTGAACTAGAAGAAGTATACTGGCTCTTAACCACCTTTGCCAAAAATTAACACTTGTCAG TCATGGTCATATTCATTTGGTCCAAATCAATCATATCGTATCAACCTAACTACAAAGGGG ${\tt ATTGGGAGATGGTGATGTCTCTGTCACAGAATCTATATAATAGTTAAAAGTATTTTTAAC}$ TTGCATAGACTCAGAACAAGATAATTTGGAGGAATTCAATGCTTAATGGCATACCACTAA GATAAGCTGATAGATATATCGTTGCGATTTGGGTCTCTGACAATAGAGGCAATTGATAAT ATTAAGAGACTATGTGCCAATTATTGTGCTTGGATTGAGGGTACAAAGGTAATAGAATCC AAGGAACCTGCACTCTTTTTGAAAGATAGACACATAAACACATACTTTTAAAATAACGTG GTAAGTGCTACTATGACAGATGGTTGCACAGAATGTAGTGGAAGTATTTGAGAAGGACAC

FIG. 1J

TTAGCTCTGCTGGGGGATTAGAGAGAGATACAGGAGGAGATGACACCTAAACTGAGTTTT AATAGATGAATTCAAGTTACCCAGGTGAAGAAATTGGGTAAGGATGTTCTAAGCAGAGG AAACAACATAAGCAAAATCAAAGAGGCGTGAAATAGAATGAGCTATGAAGAAAGTGTTAG GCAATTGGGTAAGTCCAATGTAAGTGCAGATGAGGAGGTCTGGAAATGAGGCTGAAGCA GTAAATAAGGATTGGCCATAAAAGACCTTGTGTACAATTCTTAAGATCTAGGCTTTGACA CTGTTGTTTAGGGGGGGGCTGTTAAAGGATTTTAAATTAGAGTACCATCATTGGTTTGCAT GAAGGTTTTCACACTGGGGTTTGCATCCTGTTTTGGCAATAAGCTTGTTTTAATGAAAAC AAACAAACAAACTGACAATAAAGAACATAATCCAAATTCTCCAGATAATTACTTCCAGGA GGCTTTCTACGTGCTGCATACAAAACAAAGAAAGAAAAACATAAAGTGAGAAAACGAAGG AAAAACAAGGAAAGAAGAAGAATACATATTGGAAAAACTGTTGCTGTTTTTGT GATAAAGTCTCACTCTGTTGCCCAGGCTGGAGTGCAGTGGCGCCATTTCAGCTCACTGCA ACCTCCGCCTTCCAGGTCCCAGTGATTCTCCTGCCTCAGCCTCCCCAGTAGCTGGGACTT CAGACATGCACCATCACGAGCAGCTAATTTTTTGAATTCTTAGTAGAGATGGGATTTCAC CGTGTTGCTCAGACTGATCTTTAACTCCTGAGCACAGGCAATCCGCCCACCTTGGCCTCC CAAAGTGCTAGGATTACAGGCGAGAGCCACTGCACCCAGGCGCAGGTTTTCTTTATGATG TTTTAATTATATCTTTCTTGGAACATATATGTATGAATCTTGCATGCCATAGGTCTATTA ATATTTTCCAATATTCTACATGGTTTTTTACTAAAATCATTTTTATGATTAGTTACTGAC TGAGGTTTCAATGCATCACTGTACTCCTAGCTATCTCTCATTTTAGCTTTTACATCACAT TTTGGCCTCACACTGAAACACAAAATATTAAAAATTTGAGATCTAATAAACAATTTTCAC ATTTTCCAACTAAATCCCCACTTCTTTCTAAATTTTCTACAACTTTCTAAACATTCTCAC GACCCCAAGTGAGCCCTTAGGGAATTTCCGTGAATATTTCCCTACAGGTTGGCATGGTAA CACACTTCACAATTTCTAAATCTGTGGATAGTTTAGAAGCTTTTATTTGCTGTTCCTAGT TCACAATGGAAATACAACAATGATTAAAAATTATAATATCCTTTTGTAGATTCTTAGCTT TTATTCCTACTCAGTGACTCTAAAATGAATTTATAAGGCCCATGGTTTATAACCATGTGA GGCCTTGATTTTGTCACTACATTGCTAGAAATGGGGTCAGAAGGCCACCAGCTTTAATAA TTTAATTCATCAATTCGGAATGAATTTGATGAGTCAACCACTTTGGTAGAGAACCATATT GCTCATAAATACTGTTTTGAAGGCAATTCGTCTTTCATAAAATGTGAAGATTGTGCTGAT $\tt CTTTCTGGGCAGGGTTATGGAGGTGTGATTAAATGCTTAAGAAACCATTTTGTTATTATA$ TTAAACCGAATCAACTTTTTATTATTAAAAAATAGATAAAAACTTAGCATCCTCAATTATA ATACTTTATACAAAAGTTTCCCAATTTTATATAGACTGAAGATAAAAATACATTAACAAA TCTTACCAGCTGGTTCAGGAAAATAACTTCATAATTATTGAGACATTTATGTGTTTTGGGC TTGATTTATACTTTGGACACAGGAAAACCTAGAGAGATCTGGTTCTTTGAAATCATCAGA GATGGTGATGGTGACTCAGAGATTCCTGAAAATCAGTAAGATTACCCTAGTTTATAGACG TATGTGTTATTTTTCCCCCAGGCATAATGAACTTTATAACTTGTCATTGACAAGAAGCC AAACATGTACATACCTCACACATGTGTACACACACAGTTTGGGGATTGGATGATATGAAT AATATAATTAATACACCCTAATTTTTCATGCAGGATTAAGAAAGTATCTTCCAAACATTA AAAATGCTGAAAACTGGACATAAGGCCTTGAGTTTCCCAAATTCAGGACATATTTTCAAC TATCCCCTGAGTAAATGAACTATAACATTTACAGAAGTAAAAATGATAAATACACTAAAG CTGTAGCATGATTTCTTTTCCTTGAATAGACAATATTCCTTGACAATCTTTCTGTAAACA GAATACAATGTTTCCCTAAGCAATATATGCGTGCTCTAGAGTTTTCACAATTTCTGATCC TCCTATGACTGGCTCCTGCTCAGCTCACACTGCACTTTCATGGAAGTTCTCTTAGAATGC CAGCTTTGAATCACTGCTCCCTCATGTGCTGTGTGTGATAGCATCCCATTTTAGTTTTGT CATAGAATTGATTACCATTTCAAATTGAATTGTTAATTTATTGTTCATTTTTCTGTTGTC AAGTGTAAGACTTCCAGCAGGAGGAATTTTTTACATATAAGTACATTTTTTAAATTAAGC ATTGCAGGCTTTAAATTTCTTCTATATAAATATTTAAAATAAAGCTTCAATAATTTGAAT TGCTTTTGTGATTATTTTGTTTTATACCTTGAGTAACTTATACATCAACTATTTTGTAGT TATTCTAGTAATGATATGAAAGACCATTTGAAAATCTTTCCCCAGCACTGAGATCTCCT TGACATGACTAAGTGATTTATACTATGCAATTATATTGCTCTTCTCAAGAAAAGCAAAAT GAAATTTACAAATTTGGTAGCTTTTTGTTCTTTTGTTTTCTCAAGTAAGATACACCAAGA TTTCTTTAAATGATACGCTATATTTCTGCAATAACTGAGAAGAACATGTAATGTGCAAAA

FIG. 1K

CTCTTAAACTCTTTTTGTTTCAAAATAATTCTTGGTTGTTTTTATAAAAGTCTAAGCAAA TACTTAATGAACTGTGTCCCAAATGAGGTGAAACAGCTGTGACAGAATGTTACTATGACT CTGTACTTTCTATAATAAAAAGGGACAGACATATCCTCACCTGAGCCTTGGGATGTTTCA GGCATGCCCATAGAGCCTAAGCTTTAGGAATCCTCTGTCATTCTTTTCCATTGCCAGTGA CTTGTGCCAATTCTAGGGTTCTGGACTGTGCAAACAATGGAAAAAATAATAACACTTTCA AGAATAGGAGAAACACTAATCCCATCTAATTCTGCCTTCAAACTCCTAAAATATTCATCA CAGGTCTGGTGGCTCATGCCTGTAATCCCAGCACTTTGGGATGCTGAGGCGGGTGGATCA TGAGTTCAAGAGATCGAGACCATCCTGGCCAACATGGTAAAACCCCATCTCTACTAAAAA ACAAACAACAAAAAATTAGCTGGGCTTGGTGGCATGCGCCTGTAGTCCCAGCTACTTGG GAAGCTGAGGCAGGAGATCACTTCAACCCGGGAGACGGAGGTTGCAGTGAGCCAAGATG AGCAGATCCTGGAACAACTGAACCAAATTTATTAATATGTATTATTACTGAAAATCAGTA ATGAACAAAATTTACAGAATGGGCTTCTTGGAGTTGTTACATTTCCCTTATTACATAACT CTTCAATAAAAGTGTTTGTCATACCTATTTTAGTTAATTCTACAACAACTAGTGTGATAG GGCTATTATTTGATCTTTTTTTTTTTTTTTTTTTTACAGGTAGTGACATTCAGTATTA GACAGCTGCTATTGTGTTAGTTGTCTGAATACCTTTACATATTATCAACTGGCCTTTTCA TTCCTGAGTTGTGAGTAAATGCTCTGTCTCCCAGACTGGAGTGCAGTGGCGCAATCTCGC CTCAGTGCAAGCTCCGGCCTCCCGGGTTCACACCATTCTCCAGCCTCAGCCTCCCGAGTAG GAGACAGGTTTTCACCATGTTAGCCAGGATGGTCTTGATCTCCTGACCTCGTGATCCACC CGCCTCAGCCTCCCAAAGTGCTGGGATTACAGGCATGAGCCACCACACCCGGCCATAAAT GCAGTCTTGTGTTCCCCACTTCCATTCCTCCTTTGACAGTACAGCTATGCTAGTCTGCGT AGCAAATTGAAAAAATATGACCTGTGGGATTTAAACAAAACACAGTGTCATACACATTTT CTGGTAAACTTAACCAAAAGGGACTTGGGTTCCATAACTAATCACCAATGCCTCAGTGAT CTGTAACTCCTTGTAGGTACCTGATCACAGTTACTAAAGGGAAAGAGGGGGGGAATAC AAGAGCAAAGTCAAGCCAGACATAGATTTTATCTCTTTGTAAACAGGAGTTCAGAAGACC GCTCTGAATGCTGAGTTAGCATCAGCAATAATAGAAATATATGCAGATTGTTGATTTGAA GTCATGCAAAGATATCTTTTTCATCCAAATGGAGGCAAAAGCATCATAGAGCACCAGAGG GCTAAATCCAACTGTAGCAGCAAAAGGTACACAGAAAAATAAAGCATCCTGAACCAACGC ATATTTATTGAACACCCACAATGTGTAATCTGTTCTATTACATTCTGTGGAGGAAATACA GAAGTGAATGAGGCATGGTTCTTACCTACAAGGAATTTCTAATCTTGTGGGGGAGACTAA CATGTAAACAATAAACTATAGTATGAGGATTACTGAAGAGGCATATGCTAAGTCTCAGAA TAGCATTTTGAGTTATTGTTGACATGTGAATACGATTTTGAAAAGTTCCAAAGAATGAAA AATTCCACCTACATTGGTGAAGTACTAAGATTAAATGCATGATAGCTTGAAGACACAAAA ATAATTATTATAAACCATTCCAAAAATCATTCAGGGAATTCCAATAATACACAAGTTTT TAAACACATTTCTGGGTAATTTTGAGTAATAAGGTCTTAATCTCCTCTACTGCTTTCAAT TGTTTTTGTGGCCTTCTTTATTTTGTGGGTATCTGGCCCAGTCTTGTCTGTAGTGTATTA TGGTGGATTGGATTAAACATGTTTTGCAATCTCTGGAGTGATTTTAAAATGACTTGTGTT ATATCAGAGTTTCCTAAAGGGAGATTAATTTGGCTTAATGGTAAGAACGGATTAAAGTTA TGAGATACCAGACACTGGGAAAACAGTTAGAAGCCTGTTGAGACTCTTCAGGGCAGTTG**T** GGCAGTCTGAAGATGGAAGTTGGCAACTCATCAAATGTGAGAAATTTATAGGAACAGAAA AGAACCTGCTGATTAATATAAATTTTCTGCCAAAGAAAGTACAGTGGCTCTCCTCAGCAA ACTAACATGGGAACATAAAACTAAACACTGCATGTTCTCACTTATAAGCAGAGCTGAACA ATGGGAACACATGGACACAGGGAGTGGGACATCACACACTGGGGCCTGTTGTCGGGACTA TGGGAGGGAGACCATCAGGATAAATAGCTAAAGCATGTGGGACTTAATACCTAGGTGATG GGTTGATAGGTGTAGCAAACTATGATGACACACGTTTACCTATGTAACAAACCTGCACGT GTATAGAAGGGTCATGATGAATTGGACAAAGATACGTGGAGTTTGAATTGCTAGAGGAGT ACCCACGTGCAGTTGTCCAGCAGAAATCAGGGCTTGTTCCCCAACATGCTATTCACAATC

FIG. 1L

ATGTGAAAATGTTTTAGAAAATGTCTTCTGGAATAATTAAAAGCATACAAGGGAATGTAA ATCTCTTAATGTGACAAGACCTTTTTGCCACAATAAACAAATTCATTAGTTCAAAAAATA TTTATTGTGTGCCTATTGCAGCAAACAAAACAGACGAAGCTCCTTCTTGTAGGGAACTTA TACTCTAGTGATATTTAGTATATTTTTGACAATTGAACCAACAGGATTTGCTGACGGAT AGGTGCCATTTACTGAGATACAGGGCCGTAGAGGAGGAGTGTGTTTGCAGCAGGGAAGGA GAAGACTCAAAATTTGGTTTTGATCATACTAAATTTGATATAGTACAGGTAAGTGTATGG TGGCCATTAGAACATGAAGGTAAGAGTTTAGATAAGGAGACAGGTATGGTGAAATACATC CAATATTTATAACCAATATTATCTTTTGTGTCTGTACCTTTTTATACATTCCCCATATAT ATCAAAGACTATAGAAGGGACTGGATAGTGAATAAGTGATTATACATAAATTCTTTTTTA CAGATTATTTTGCTCTTGATTTCTCCTATGTAAATCATCACAGCTACATTTTTTAAAATC TTAAAAAGGATTACTTTGAACAATGCATTTAAACATCCAGAAAACAAAAACAGGAGTGCA TGGTAAAAATTCTGATTTCAGAACGTATGCCTGACTTATCAAGTCAGAATTTCAGGGAGT GAAGACCCTGGAATCTACACTTTAAATAGAGCCTCAGTTCACCAAGTATGAGAAGTCCTG TAACAGGGAAAAGTAACCTCCTGTTATATTTGATGGAGGCCAATTGACAAGCCAAGTAGT TTTCCATTTGACAAAAATTCTATTGTACCAATGAAGAGCTATCAGAGGGGAGTAGATTAA AACACCTCCCTTGAAATGGAATTTGGCAAGAAAGCAAGAAATTACAGCAAAAAGACCAAT AAGAGGAATTAGGGGCAATGAAGGAAGGAGCAAAGATGTGGGGAACCCAAAAAGTTTTCCT TACATTGGTGTAATTTATTATTATTAAGCCAACAATATACTTTTAAACTTATACAACT TTGCAAAAAGTACAAATCAGAAGTCTGGGCTAAGTAGAATGCATAATAGAATCAGTAGT ATTAAAATGAATGTCACTTCTTTTTACCATGTGTCCTTTAAATTATTAAAATCTATACAC ATATTGCTATACATAGTAAATATAGTTAGTCAATTATGTCATGGAAAGAATTGAAGGGTT GTTATAAATTTAAAGGTGTTTCACTATACAAAAACATTGTGAAATACTGGTGCTGATTTA GTTCTAGTATCTCTGATATATTAAATCATAAATGTCAGGAGTTATTGGTCACAAAATAAA CACCAGAATTATATGACAGTCTAAAAACAAAAACAAAAACTTCAGCAACAATATTGAAG ATATGGAAGTGCCAGAAGAATAAGGATTAAGACAATGAATAAAAATCTCTTCCAAGGACT GGTCTACACTAAGAGTTTAGAAATGCATTTTTTTTTCACAGAAATATCCTTAATCCTCTA TATAGAAATGAGAAGAAAACATAAGACTTTAGCAAGCTCCATCTAATCCATTTGCAGACA TATGGTTACCTATCTTTCTTCAATATATTGGAGTTTGCAAATATTCTACCTTCAAAGAA TAGGTGTTACCAAAACATTGTCTGCAAGATTTCTAAGATTTGAAATATATTTGCTATAGT AGGTTAGAGATGAGACATTTTTACTTTAAATTGCAATAATTCAGACTTAAAATATAAAAT TGTGTATACTCATATGAACTTTAAGGAAATATCAGAGGCATCAGTAATAGATAACTTGCA TCTCTTTTACATTCAGTTCAAGCTACTCAAATTTTAATCTTTTGTTTTCATTCCAACAAA AAAAATTAGGATCTGCCTTGGCTTTTGCTAAGAAAGTAATTATTGGCTGGACATGGTGGC TCACATCTGTAATCCCAGTACTTTGGGAAGCTGAGGTGGACAGATTGCTTGAGCTCAGGA CGCGCGCACACACACACACACACACACACACAAATTAGCTGGGAATGATTACACGC CTGTGGTCCCAGATACTTGGGAGGCTGAGGTGGGAAAATCACCTGAGCCCAGGAAGTCGA TTATCTTCAACACTGTGCATACACACTTTTCTGCATCTAGATCCCAAATTTTTGTTTTGT ATTTACATAGAACATTGATAAGTAAGGTAAGTATTAATTGATAAAACATTTCAAACTCAT TTTTCACTAAATCCAATGGCCTTCCTCTTTTGCATGAAGTCTCTAAGAATCATGTTAATC TACATACTCAATCTACGTAACAACTGGATATATCCTGTAGTTGTTGCCCATTTTTCTGCT AAATGTTATCTTTAGCACTAAGCATGAGTATGAGGAAACAGTATCTGTGCTCAGATTCCA GAAATGAAGAAAATGTACTGGAGGTCTTTTGGATAATGGCTACAAGGTCACAGGGACTGA CTCCTTTGGAAGCTCAGCGATAACCATTTTCAGAGAGAATATGTCAACATCTTTCAGTCT AGAACTTGATGTTCTGCTGAGATCTAATCTGGGGGTGTCCTACTATTGAATAGGTATAAA ATATTGAAAGTCTATTAATTGGCAAGCACTCTTCTGACATTAGAAGGAGCAAAGATAAAA AAGATATTATCATTAACCTCAAGGACATGACAGCATCATGGGAAGGCCAGAAATGCAATA TGTTAAAGTAAAACACAGTGTAGTGTTTACTACTAAAGAGATATAAACAGAGTACTGTGG TCTAAAATCATATATATAACATTTGCTTAATGGATGAGAAGGAAACTTTAACTTCAGGAG GCAGAGCATTAAGAAAGTGAATGACAGGAGGGTCAAAAGAAAAAGCCGACAGTGTTGCAG AGGCAGGGCATAAAGGAGCTAAACCTTTGCTACCTTCAGTTTTTATTATCCACAGAACGA

FIG. 1M

14/51

AATTTATGGGGGCATTCCTATGGTCCTCACCTCACCCCATTTTTCTGTTTTACCTATGAA **ACTTGATCAAAATACTGTCTCCACATTTCTCATAAAATACATTAGTTTAATTTTCTACTA** TTACTTTCTTTTAGTTGATTTAAAAAAAGGTCATTTATGACCTATTTAGGTTAGCATCAT TAATTTTATCAATGTAAGAATATGGTAGTACAGTGTGAATTCCATTAATGGATATGTTGA TACCATGGGTTTCTCTGACCTTTCCTCTTCCGCTCCTCCTGATGATTGGTTCTGAGCTT TTTAACACTGAATAAAGATTTTTTTTTTCTCTAACAGACTTAAAAATAGTGCCCTAAAAAT AAAAGAAAAAACAAAAAACAAAAATAAACATTGTGTCCTACATTTGTATTAACTTTCTTA TTGAGTTTTATTGATAATACCTTTAGGATGCATGTATTATTAGAAACATCAGTTATTTAC **AAGTTCACCTATTTAAAAGTCTAATAGGAAAAAATATTTCATGTTGCTAAGTATGTGACT** TCCCTTTAAAAGATAATAATGCTTTCCCTTTAAACAACAATAGTAAAAGAAGTAGAGTTC CTTTTAAACACATACTTTTATATTATAACCCATTCTGTTTAAAAAATAGCAGGCATATAA TCTAGAAATGCAAATAATTTAGTGAAATTTTTAAAATTATTCTACATATAATTAAATATG GATATTCGTTTTCAAATATCAAATAATAAAATATGTCTGAGATGCTGACTAATCCTTAAT TATAGGTGTGATTTCTACTTCACCATCAATACTATGGTACTCCAAATCTTAACATGAGTC TGATTTTCTAATAAACATGATGAAAAAAGTTATGGAAAAATTTTGAGATTTACTTTGGGA GGTTCTATTGTGTTCTGTTCAGCTTCATAATATTCAGTTTCTATGAGTTTTGGTATTTAAT TATGTGTGTTTGTCATTCAGTAGGCTGGAAGTATGACCATTGGGAAGATCAAAACGATAAG ACATTAATGACAGTGCTTTATCACTGAATCTAGTACTTTTTTTAATGAAAGAGATGTTGG CCTCTTGTATTGTTATAAAACAACACAATTTTATGGCTTTAAATTAAAGTACAATCATAA CAGAAGACAAATTAGATTAAAAAACAAACATGGAGTGACTCATATAAAATATTTAGAAA CCAATAATACAGATAGAGACACATTAGTTCCTCTAGACATTGTGTTTTTCCAGTAAAATGA TCACCAAACTTACCAGGAAAATGATAATTATCAGATTATTTACTTTCAGAATTAAAGGCA GGAAGAGAAAAAATGAATGAAGAGGAAACACAGTAACCATATAGGACAATAAGAGTGAA TGAAGATAAAATGAAAAATCAATAAGATATCGACTTTCTTAAAAGACAAATATCACAATA GGAAACACCTCAGAAAGGGAAATCTCAAGAAAATAATAAACTGAAAGAAGAAAACATATC AAAACAACTTGAGGACTGACAAAGTTTTAAAATGTATTTAGATAAAGATACCATGAGGAA AGTGATCAAGGTGTTCTAGGTAATCACTGAAGATAAAACTAAAAATAGCTTAAATTAAAA TGAACATCTTAACAATGTCTCAAATGTCAGGAATTGATCCAGTTTTTTGGCTGCACAACAG AGTGGCTATAGTTAACAATAATTCACTGTATATTTCAAAATAACTCAAAGAGTAGAATCG GAATGTTGCTAACACAAAGAAATGATAAATTCTTGAGGAAATGGATATCCCAATTACCCT GATTTGATCTTTACACATTGTATGCTTATATAAAAACAGTATTCATGGCCGGGCGTGGTG GCTCACACCTGTAATCCCTGCACTTTGGGAGGTCGAGGTGGGCGGATCACAAGATCAGGA GATTGAGACCATCCTGTGAATGGTGAAACCCCGTCTCTACTAAAAATACAAAAATTAGC CGGGTGTGGTGGGCGCCTGTAGTCCCAGCTACTGGGGAGGCTGAGGTGGGAGAATGG CATGAACCCAGGAGGCAGAGCTTGCTTGCAGTGAGCTGTGATTGCACCACTGCACTCCAG CCTGGGCGACAGAGCGAGACTTCGTCTCAATAAAACAACAACAACAACAACAACAAAAAAC ATTTAAATTTTTCCTATAGCATAGAGATCTGTAATTAATACTTGTCGATCATTGTTGTTT CTGTCTTCCCAACAACTACACTCCTGTTTCTTCACATTCCCCCTTCTTCTAACAGCACTA CATCTTTCTTTAGGAAACTATCCTTTTGCCATTTCATGTATATGGTGGGGTGGGGGGAGTT ATCAATCACAGTACCCCAGCAGATGGGACCAGAGGCAAAAATGCCTGACCTTCTCCCATC CCCCAACCACAGCAGCAAATGAATTATAATTTGATGCACAAGGAAGTATCGGAGCTTTTG TGTTGGGTTTTACATATCACCTGTGGGAGATAAATGAACTTTTCCCCACCTAACCTTTAG CCACTTGGGATGATTAGACATAGAGGTGCCTAAGATCTTTCCCTTTGCCACATTAAAAAC AAATCATCTATGGCACGAGCATACAAGACCAGCTTTCAGAGACACAAAATGATGGAGAGA ACCATGATACTAGTTTTAGACCTAGTCACTGAGACTTTCTCTGCTCCTTCCCAGTTACCT GAGCTTTATTTTGTTTACATTTATCAGATTTGAATGGCTGTACTTCAAAGTACTGATTAA TTAACAAAAGAGAATAAAGAAAAAAAAGAAAGAAAGAAACAAGAAACTCTGACTACCTCTCC TCTTTGACATAGTTTACACTTCTGACAGATTGTTCTTCTCTAAATTTATGTAGAGATTAG AGTGAGGATGATGTATGCACTGTAGCATGGGTGGTCTTCCAGGAAGCCTTGACTGAATGA

FIG. 1N

15/51

GGCAAGGAGTATGTTGCTCCCTCAGTAACCTCAAATTTACCTGCAAGCCTGATAAAAATC TAACACTAACACTAAACCCAATCTTATCTACAGCCCTAACTGCACCCTAATATTAACAAC CCTACCTCTGTACTTCAAAACTAAAACTAATTCTGATTTTACTCCCATCTGCCCCTTTTA CCCTAAAACCAACTGTAAAACTAAATTTAACTCTAAACGTAATCCTAAAACTAAGAATTA ACTAACAATTTTATCTCTATACCCAACTGTTAACCCCAAGCCTAACTCTAATCCTATCTC TAACCTAACATTAACCACAAACCTACTTCTAACTCTAATCCTAACCATAACCTCAAATCT AACTCTGATTCCAATTGTAATCTAAACACCAACCCACCCTACCCTTTTATCCCAAATCC ATCTGAAACCCTCATCAGAACACAAATTCCAATTCTACTTCCCACCCTGACTCTGACTCT AAACATAGGCCCAAATATAACTCTAACTCGAAGTCAAAAACTTAACAAACCTTATCTTGA AACTCAACCTTTACACTAACCCCAATTCTATCTGTAATCCTAACCCTAATATTATCATCA AACCTATGTCTAACACGACCTCCAACCCAAAACCAAAACTAACCTCAGACCTAACTCTAC ATCTAATTATAACCCAAACCCCAGGCTGCTACTTACCATAACCCTGAAACTAAGCTTGAT CCTTTCTCTTTTTTTTGAGATGGAGTCTCGCTCTGTCTCCCAGGTTGAAGTGCAGTGGCG TGATCTCGGCTCACTGCAAGCTCTGCCTCTCAGGTTCATGCCATTCTCCTGCCTCAGCCT CCCGAGTAGCTGGGACTACAGGTGCCCGCCACCATGCCTGGCTAATTTGTTGTATTTTTG GCAGAGATGGGGTTTCACCCTGTTAGCAAGGATGGTCTCAATCCCCTGACCTTGTGATCT GCCTGCCTCGGCCTCCCAAAGTGCTGGGATTACAGGCATGAGCCACCACGCCCAGCCGAC CCTTTCTCTTAACCTACACTAACACTAACTGTAAACCTAGCTGTAACTCTAATTGTAAAC CTAACCTGATTACTCACTACAAAGGTCCCTCTAATTCTAACAAGAAACTCAATCCTATCT CAATTCCACCCAACCCCAAAAGTAAATCTAAACTTAAACATAACTCAAAATGTATCTCAA ACCTTAACCTTCACGAAACTACATGTAGCACTAATGTAACCCTAAGCCCAATCCTATCAG TAACACTAATGCTAAAACAAACCCCAATCTGTATCTCTACCCCATCACTGACACTACCCA ATATCTCCAACCCTAACCCTAACATTAACCCCAAACTTATCATTAATCATACATCTAGCT CTAAACCTAACCCCAACTTTAACTCTTACCCTAGCTCTAAAATTAACCCCAACACTATCT CAAACTGTAATCCTAATGCTAACGTTCAGGCTACTTTTAACCCTCACCCTACACAAAATC CTGCACCTAAACTCAACCCTAACTTTAAACCTACCTCTAATCCAAACACTAAATTTAAAT CTGGATCACGACTTGGGCATACTAGCACCCACTAGTGTTCTGGGTGTCATTTCTTTGCTT CACTCTCATCCAGCTTTCTTTACTAATTTTGGATAATGAATCAGAAAATAGTGGTGTGGA ATCAGGGTCTTTGAATTATTGTATTATCCAGAGTTTGTCCTGCTGCAAATGATAAAACAC TTATTGGCTCAAGGAACAGAAGAATTCAGTACTGAGTTTCACAAATACCTGGATTCCCTA TTGGATTCATTGTTAGGTTTCCTGTGGCAAGATGAAATGGCCTCAGGCCTGTAACACAAT AGGATCAATTACAACAGAAGATAGTATTTCTGTTTTCCTGGTTGCTCAAGCCTAAATTCC AAGATTAGTTTATATCAAACCTAGTTAGTTTTGCTCATGTAAGGGATTACTGCAACTGGG TACACTAATATGAAGAGTGGGAGAGTTGGTTAAGGGGGTTCTCTGAAAGGAGAATTAGGT CAAATGTTGAGGAAGAATTTTCCTTTATGATCATCTAGCCCAACTTTTTAATTTCTAATT TTGTGGTTTTGACCAGTTTTTTGTTTTTTTTTTAATGCAGCATGTCATAAAGTTGGGAA TACTTCACATTTTGTCTTTGAAAATTTGGAGAGTACTTAAAAAGATTTACAAAGGGGAGG ATTGAATTATTTTAGGAATGTAAATGGTGGTCTTCTGTCTCAGGCAATGTAGATGCTTGC TAGAAAACAGCTGACTCATGACTGTTTTCTTTCTAATTCATTAATATGAATTATTTCAAA CTGCAAAGTTATCTCCTTTCTTCTCCTAATCTATCCACTTAGAGTATACATGTTCAAATT AATGTATTGAACTAATTTTTCTAGTAATACATTCTATGCATTACAAAAATAGCAGTGGGA AGGTGAAAACAAATGCAGTTATGCATTTATCTCTAAATGTGTTCAACATCTCTTATGCG TACTTCAAAATAATTACATTTGTTTAATTTTGAAAAAAATATTAACAAGAAGTTGTAATT TGGGGAAAATTTAAAGCTGGCGAAAAAGGCTTCATCATAATTGACAATATGGGAAAATAC TGTATTAAAATCCTAGGTTTCTCCCTTGTTTGCATGAAGGAAATGAAAAATATATAAGGG AAGGATTTAATCAGTCAGGCAAAAATCTAAATTCATCACAGGTTTATTCACTGCATACTA TAAATGAGCAGAGTGTGGTGATTAAAGGTTGGTATTTGGTGCTGGGTAGACCCAGCTTTG CCACTTACTGCCCAAGTAAATATTGCCATCCATCAGATATCTCCACCTATCAGACCCACC CTGTTGTAATAACAAGATTAAAATCTGTATCACTAAAACTTTAAAAGAATTTATAGCCGA TGTGTGTGTTTTTGAAACGGAGTCTTGCTCTGTCGCCTAGGCTGGAGTGCAATGGTGGGA

FIG. 10

16/51

TCTTGGCTCACTGCAACTCCCGCCTCCTGGGTTCACGTGATTCTCCTGCCTCAGCCTCCT GAGCAACTGGGAGTACAGGCGCACACCACCACCCAGCTAATTTTTTGTATTTTAGTA GAGATGGGGTTTCACTATGTTGGCCAGACTGGTGTTTTTTGAAAAGACTTTTTCCTGATT CAGAAGGTGGGACTCACAATTGTAATTCTGCTAATGGTTGTCTTTCAGTCTATCAATTGC TTCATAAATGCATCCACTGTTCCTTCTTCTTCTGCCCTGCTTATAATTTTCCATGAGTCC ATATATCTTTTTACACTGTCTTTAGTCTTATTCACTAAATTAAAAACTAATTTTTGATAT TTGGTATTCATGACAAGACAATTAGTAGAATTTTGATGCTTCTTGTCTGCAATTACAGAA TCAATATTTTCTATATTATTGTATATTCTCTAAATCTTATTTTGTATAATAGCTTTCA GCATGTTCTTTAATTCTGTTTAGATATTTAGAAAGTATTTGTTGTTATTCTGTAATTTAT TTCAATATTCAATTATAGTTTAATATTTTGTTATCTAGTGTTGTCTTGATTTTGATATAC GTACTGATTTTGTAGATCCAAATTCCTCTTTCCTATCAGAGAATGCAATTTTTTACTTGG ATAAATAAGAATCATATCTCCTCTGCTTGCTACCGTATTGCATACATTCATGGGTAGAGA AAGAGTTAAGCTGATGAGAGTAGGAATTAAGGTAGACCTGTTTGGTAGGTTCTCCCAGAT TTCAGAGGACAGACATCTTTTTTTCCCTGCCTTGGTCATTTAAACTTTTTGGATTTTGGA TTCTCTGCTTTAGGCAGAGAAGCATATGTAGTCCAAGAATGTGCTTTTCTATCCAGCTAC ATCAATAATAACAATTAGTAAAATTCTACTTAAACTTAGACCTTTGCTGTTCTCTTTTCT CTGCTTGTGTTAAGTCATGCTCATGATTCTGGCAGTTTTCCACAGTACCATGTACAGAAA GCTTGAATAAGGTACATCTAGAATACTCATATATGTTCACTTCAAAAACACATTTTTGTG GAATTCTAAATGCAAATCTCAATAGTGCAATTCTAATTTACAATGAGAAAAAACTAAGGG ATTTTTTCTGGTGATTCTTTTTGCTCATTTATAAATATGTTTTTAAATGGTAAGCAAATA TATAAATTAAGCTTTTCCTTACGTAGCTACATTGATTTACTAGTGGTGGAAAAGGTTAAG CAAAACTAATTTTCATGAGTGTAAATGAATTAGTAAGTGACATATGCAATGCTTAAGGGG AATTTGCATAAATCTATGACTGATACTCAACCTCTTGCTTAGCGAGAAGATAATTAAAAT ATTTTATACTTCAAGAAGACCTAGTTTTCCAAATTATTTACATCCACAAACTCAGATTTT ATAGCAAGTAAGAAAAGTTAAGTCAGAAGCATATACTATTAACAGCTACTTACATTGCTC AAATTTAATATACGATTGCTGCTTTTGTTGGTTTTGAAATGTTTCTTGACCATGGATCTG GATACAATAATATTAAAAAATTAAATGTAAGTTCCTGCACTCACAGTAGAGGTAAGTTCA AGGTTATAAGAGAGCTTATAGATTCTGAGATTTGGAAAGAAGAGAAAAAAACTTTT CAGATTAAATAATGTGTTAATTGTGCTTCTAAAACAGCTTTGGTGATCTTAATAAAATAA ATATTGTTTTTATTTCCATTTTTGCTTTTCAGACAAGAAATGCTACTTGATGGCTGCATA TATTTGTTTTGTCTCTTTTCACCACCTACTCTTGCTAAATACTCTCAACCCACTCATGAA ATTAAAGCACATTGGAAAACATTTATCAACTACCTGTAAATACAACCTATGCTCTCTTT GTGGAGGTGATAGACATTCATCAATGGAATAGTTGATCTAAATCCTAGTCTTCATTATCT TGTTTTATACATTCTTGTCTTAATCAGTTTGGGCTGCTCTAACACAATACCATAGACTAG GTGGCTGATGAACAACAGAAATTTGTTTCCGACTGTTTTGGAGACTGGGAAGTCCAAGAT CGAATTTTATGTCTGGTGAGGGCCTGTTTCCTAATTAATAAACATCTGTTGTCTCATATG TCCTCACATGATAGAAGGGGCAAAGGAGCTCTCTGATGTCTCTTTTTTAGAATATTAATC TCGTTCATGAAGGCTCTGCTCTCATGACCTATTCCTTCCCAAAGGGCCCACTTCCAAAGA CCATCATATTAGGGATTAGGTTTCAACAAATGAAGCCAGGGGGAGGTTGGTAAACATTCA ATCTATAGCAATGCCTATCTCCAGGAGCTGCCTGTGGAAACACTTTTATCTGATATGGTA GTTTAAAGCATGGCAGGGATAAGTGGTATGAGGAAAACTCTCCCTGCCACCCAACGCACA CATCCCACTTAAGCTTCAGCAGCTCCAATTTTATCTGTGTAATATTTGGTTCCACATCAA AGTTGTTTTGAATATACTTCCATTACCTTAAAAAATGTAAAAACACTGCTTTAAAAAGCC AAGCCTATTCCCTTTTCATTATTCAGAGTTCTTCCAGTTTTACCGTTACATCAAATTAGA ACTACATAATTAGGAACCCCTCTCTAAATTTGCCTCTATACAGAGAAAAACTGTGCCTGA GGACAGAATTTGTACTTCGTTCCATACATAAAAACTCATTTGACAAATAACAAGCATAGC TCCAAGCTCAAAGAATAGCTTAATTTTTCCTGATTAGTTTATATCTCTCTTATTAATCAA TGACATTTAATATTACAACCATAGCTTGGGGTTTTAGTTTATTTGCTTTCTATCTTTTTT ATACTGTCGGCCTACCTGTGCCCAACTATGTTATAGTCAGGGGTTGGTAAAATAAAGACA AAACAAATCCTGTCTTCCTGGAGATCACCTTCACTGGGGGTTGAGAAACAATAAGAACAA GTAGTAAGTAAAATATGTACATTAAAATTTTAGATGAAGTTAAGTGCTATGGAAAAAAGT AAAATGGAAGAGGTGTTATGGAGTACCTGTTCGGGTATGGGTTCAATTTACAAGTGGATG GTCACCTTCTCACTGATAAGGTGACATTTGAGCAAAAGTCTTCAGCAGGAAGGGAGAATG CCATGCAGTTATCCTAGGAAAGAACATTTCCAATATAAGTAACAGCCAGTGCAAAAGCCC

FIG. 1P

TGATGTAGATGCATACCTTAGGTATACGAGTAACAGTAAGAAATTAGTGGCACGAAAGAC AGATGTACTTGGAAACCAAAAAGAATCTCTGGTAAGAAATTGTAAGTCATTGTAAGGACT TAAGGTTTTTTTTTTCCTCCCAAATGAGATGGAGATCCATTAGAAGGGTTTGCGTAGA GAAATAATATGATCTGACTTATATTTAACAGGACTACTCTTTTGCTGAATTGAAAATTGT CTCTAAGGGTGTATATCAGATCTTATATTGATCTTACCCTTCTCTGTTCAATATTTAACA ATCCCCTCTCATAAGGGCTGTGAGGGCCTAATCTGCTTACCTATCCAGCAGGCTGGGAAT GACACAGAGCACTCACCAGGAGCACTCTCAACCTATGACTCATGGAAGTTGGTAGATGAA TACCCCAGCTCTCATATTCCTTGGGTGGAAGAGCTCTGAGATGTGTGTTCTACACCATTA CCCAGAGGGCACCCTCTGGATTAGGCTCAAGTTGCTGACAGTAGTATCTTGCTGACTAAC ATAATTTTATTAATTTTCTCCCCATTTGACCTTATTTCTCCATTTTTCTAATAGTGTTC ATTGGTATCACTTCCAAAATAAATTACCTTTACTTGAATATTTTTCTTAGAATCTTCTAT ACAAACCTGAGCTAATACTGGGGCAAAGAGTGGAAGCAGGGAAATATTTTGTAGGTGTTG TGGTGATGTAGGACAGAGCCTGATAGCTTGGATCAAGGTGGTAGCAAAGGAGATTGTAGA AGCTATCACACTCTTTATATATTTTGAAGACACAGCCAAGAGGTTTGGTGGAAAAATGGA TTGTGAGAAGTAATAAAAAGAGTGGGAGAGAAAGTCAAGGATGTCACCAAAGTTGTCCTA AGCAAGTGGAAACTTAGATTTGGGAGAATCAAAAATCCTAAAATATCCAAATCCTCTCCC CTGCCTTCCCCTCCCCTCCCCTTCCCTTTGGAGATAGGGTCTTGCTCTGTTTCAC AGGCTGTAGTTTCGCGATCTCGACTCACTGCAGCTTCGACCCCCTGGGCTGAAGT AATTCTTCTACTTTAGCCTCCCAGGCAGCTGGGACTACAGGATTGCACTAATGTGCCCAG CTCAAGCAATCCACCCTCCTCAGACTCCCAAAGTTCTGGGATTACAGGTGTGAAACACTG TGCCTGGCCCAACATTTTATTTTCAAATATTTAAGTTTTGAATGTCTATTCGATAACCAA GTAAAGAAGTCAACTAGAATATATGAGAATGGAGTTTTCTAGAGAAGTCTGGGTTGAGGA TGTACTTTTGGGAAATGGAGCACATACTTGGTATCTAAAGCTGTGAGCCGAGATGAGATC ACTAGGTAGGTAAATATAGATAAATTAGAGAAAATATCTAATAATTGAGACATGGAGTAC TATCATAAATTTTGAAAAGACAAGAAAATGTGAGAGATCGAGAAGAATGGCTGGGGAAGA AAATGCTACTGATAAGTAAAGTGAAATGTAGAATGAAAGTCAACCATAAAATTTGGCATT ATGGGGATCATTAATGACCTTAAAGAAAGTGCTTTTAGTGTAGTAATAGAAAGATGCAGA AAGTAAGTAGAGTGAATTCAAATTCAACAGAGAATAGACAGAGAGGAATTGAAGACATTT GGCAAGTGTTTGGAGGCCAATTTATACTCAAGAATAATTTCTTGAGTTGGTTTTTTTGTGT TTGTTTGTTTTTGATTGGTTAGTGTGTTTTTTTTTAGACGGGATTGGAGAAATACTTTC ATTTGTGTTTTTACCCATGTTTTCAGCCTTGCCCTGGCTGCCTGGTATAACGCAACTCTA TTTGTTATTCTGCTATTATAGTTTCCCTAGCTTGAATTTTTTTACACCCTTATTATAATT GTAGCGTTGCATGCCTATTTCAAAACATCTCATGTACCCCATAAATATATACATCTACTA TGTACCCACAAAATTAGAAATAAAAAATTTAAAAATTATGATTTTTAAAATTTGTTA TAGTTGAAGGAAATTTCAGATATTTTGGTAGCAGAAGGAACTGAGTTATGGCTCAAGAGT TTTTTAATAAGTGTGAGTGGAGTTATACAAACTACTCATTAAAATCTTTATTTGAATTTG TAATATCTGAAACCATTTTCATATTGAAGAATCACTTAAAATAGTCATAAAATGTAAAAT TGCAAGACAATTAAAAACAAAAATATGATTTCACGACTGTGATAGTACCTGAGAAATTTC TTCATCTCCTTAGTAAGAGAAGTATTACACCTATTTATAGTTATTTTATGAAACTAGCTA AGATGAATTATGTAGAAAAGATACAGATTTTCAAACAGAAACTAGAATTAATGGAAGCTA AGCGAAAAATAGCATCTACCTATAAGGATTTGCAAAGCCAGTAATCTTTCTAAAAATATC AGCAAACCCAGAATTAAGGCTTATGTTCTTAGCTCATTGTAACTAGATCAAAAATAAAGA AGGCCAAATAAAGGTATGTGACATTTGTTGAAAACCTGAAGTGTCCTATATGCAGAAATA TTTTTATCATTTAATTTCAGAAACTTCTTAACATGACATGATCCTCTTGAAAAGAT CACATCAAAAAAGGCAAAATAATTGCATAATTATTGTAGAATAATTTTTGTGTGAGTATT TTTAAAATATGCCTAATTTTCCAGGCATTGGTTTGCTTTGCTATAAAATGGGAGGATAGA AAATAACTTTCAAAATATCTTATAAATCTAAGAATCTTTGCATCTTATAAATCTAAGAAT CTTTGGAAATTCATAGATTATTGAGATGGAGTCTCGTTGCTATGCATTGTAGCAAAGTTG

FIG. 1Q

18/51

GAAATAAATTCTAAATTTTATTTCATTTATATTGATCAATAAATTGTTACATTTCACTAA TACAATAAGGAAAATTTATTTTACCTGAGTGTATGTCTAGCTTGTGAAATAAAAATGCTC AATTATGAAAGCATTTATTGCCATTTTGAATGAAAAATGTAATATGTAGAACAGAATTTT TTTTGCCTTGAACTCAGTTAAATGTAGAAATTGATAAGGACTTGCATTTTCATGAACTTA ATAATTATCTGTCTTTTCAATGGTCTCCATATCAAGTCTGAGAAATATGGATGTGATTTA TTTTAAACCTCACCATTTGAAGTAAATCTAAAGATTCCATTAGGTTATGAGCATATAGGA TACAAGGACCATATTGACAGTTTTGTGGGATTGTATTAGGATAAAAGGGTAGGAACAATG GGGAGAAATTATAGCTTACAATAGGGAAGAACCAAAAATTGTTGCAAAATGATGGAACA GGCTGAAAGAATGATATAACCTCCTAAACACTTCAAATGTTTAAGCAGTTCATTGTACCA GTTTCTTCTAACGATAAAATAGCACAACTCACTTTTTTTCTAACCTCTAAGAGTATATTA TTAAAAGTCTGCTACAACTACAGATAGAGGAACAGTTTGTAGTATCCGTGATCCTAGAAC CTGAGCTTTAATGGCATCATCATGTGATATGACTTGAGATTTATATTTGGAAGAGCTTTG AAAAATCACGGATTGTTACCCTAATGAGGTGTTATTCAGTCTTTTAAACAAGAGCAATTT ACTTGGCTGGGCATGGTGGCTCACACCTGTAATCCCAGCACTTCGGGAGGCAGAGGCTGG TGGATCACTTGAGGTCAGGAGTTTCAGACCAGCCTGGCCCAACACGGTGAAAAACAGTCT GTAATGCCAGCTACTCGGGAGGCTGAGGTGAGAATCACTTGAACCTGGGAGGTGGAGG TTGCAGTGAGCCAAGATTACACCATTGCACTCCAGTCTGGGTGACAGAGCGAGACTCCAC GATTAAATTTAATCCTGCAAGATTATGTCTTTTGATGGAAATGAGAGGGTTTATACAAAG TTTTATTCGTGATGTTATCTATGTCATCTATTGATTTCTGCTCTGATTCATGTGGATGAA GTTACACCTCACACTTTAAGCTGGTGTCAGTCTTCCCATTTTCTGCTGTGATGTGTACTC AAGATCTCCAGATTACATCTGTAATGTAATGCAGCCATGATTGTTTATAGGTACATTTAG ATGAATTCAATGATGAGTTATGTTGTAATAAGTGTCAGATTTAGATGAACCATACAAATA AAAGAACCATGCATTAAAATGACAAATGTGTAAAAGCATTATTTGGGCCTTAAGTCAAGG CCCAAATGTGGATACTGGTACTGAGACATCTTTCAGAAAGGAGGTATGAAGTACTGAAAA ATATTTACAAAATGAAGACTACTTTTATCTTACTTATCATGATTCTTTATTACATATGC ATTTTCTAAGATAACTATAGTGCATTAGTTTGTACTATGTTAATAATAATAATAGGGTAAA TCAAACAATGTTTTCTAAATCCATTAAAATAGAGTTCCCTAAGGGAGTTAAAACAATTAC GTTCTACTGTATATTATTGGCATGCTTCAGGAGACATGATTTAATCTCTAGACTATCAGA ATTCAAGAACTAGTGAGTCATATAACAAAGGAGGCTTAATCATGCCATTTAAGTGTCATG GAAAAAGGTTTATTGGTCAGGAAAAATTAATTAGAAAAAAGTTATAAAATACTTCACTAA GAAAATAAAATGTCAGGAAGCCCACTTAGACAATGAGTGAAAATGAAACAAATTCAAGTT TTTACAATATTTGGTTTCTATAGGATTGCTTCATTGTTTTTGGTTTTTTCCCCATA TTGTTCACATGCCACACAGACAATCAATTATGAAGAAAGGAGAGACTCGTAGGAGGCAGG GCCAGGCTGTTCACACTTTTAAACTAGGTAGCCACAAATGAGGCTTAGTTACAAAAACTT GAAAACTGGATTCTTCCCAATGTATTATACATCCCCAAAGAAATGATGAAGTTCCTTACT TAGGTTTTCATATTGGCTTAGATTTTTTTTTTTCATTAACTTGCAATTTGTGGTTGGGAAAT GATCTGCTTTTTGTTTCAGGTTGTTTAATGTTTTCCAATGTAATATTCTTCTTGCACTCC AGTGAGTTTATTTACAAAACATTTAATGTCATTTGCGTCTTCGAAGAACAATGTATTCGG TTAGAACAAAAGTGAGCTCCTGCATAGAGCTTATGATGGTTTATAATTGGTAAATTATTA CTCATTCAAAGGGACCGTTCACCCACAAAATGCCTTTTTGTTTATCTTTTGGAATGACAC CATTGGAAACTCAGTATGGCCACTTTTATGGTAATAAAAAGTCATATATAAAAAGGAT CTTTAAACTACTAAATACAAATAAATTAGTAGTACAGTCATTAGGATTGCTCTTAGTTTG TTAGTGTTGGAATAGACTTTTGGATTTTCTTCCTAGCTTAGATTGATACAATGTGATGGG GACTTGCTCTCCAAACACAGGAATAGGTGGCCTGCAGACACACTCTGTGATGCTGTAATT CTAATCCTCACTGAATATATCAGGGGTGGACATCTGGCCTGGGGCAATTCAGATACTTTT TCTTAAAATTTATACTACAAATTCAAAAGTGGTAACTCATCTCTGCCATCACTTATAGTA

FIG. 1R

TGGAGAATAAAATAAGTAAATTAGAACAGGAAAAATGCCAAAACACACAGACATGACCCT GATAGTTTTCCATTTCCTGATCACTGTCCCTTCCTGTGGCTGGATAAGGAACTGTCTCTA GGCTCTGTAAGACATATTTGCATCCTTACGACAAATTTCTACTCCTTTTCATAAACTAGA CTTGGGTTCTTTAACTTGCAACAGCAACAACAATAAACGATTTTGTTGGGTACAATCTGA TTTTATTAACTTCTGGATTTAAAAGCCCTTCTAAATGTTGATTGGCATTGTTTTTACTTC CTAAGAGTACGCTCATGCACCACATAGTGATGTTTTGGTCAACGACAGACTGCATTTACG ACTGTGGTCCCATAAGATTATAATACCATGCTTTTCTGTACTTTTCTATGTTTAGATATG TTCAGATACACAAATGCTTATCATTGTGTTATAATTGCCTACAGTGTTCAGTACAGTTAC ATGCTGTACAGGTTTATAGCCTAGGAGCAATTGGCTATACCCTATAGCCTAGGTGTGTAG TAGGCTATACCATTAGATTTGTGTAAGCATACCCTATGATGTTTGCACAATGATGAAATC AGAAGTACAATAACTTTCAAATCCTGAATGTTCTGTACTTTCCATCTCACAAGCATTTTG CAAAGCATCAAATGGTATAAGCCAGATTACTGTTAAGGCAACTTGGAATTAATATGCTGC TCAGTTCTGGAAAAGGCATATTCTGTAAATATAGATGAGAGAATATAGACTTTTTCCCTC TCTTCTTACAATCCACATTCTATTCAGTATTTCATTTACTTGAGGGGTTATATGCTACTT ATCTTTATCTGTTGTGGAGTGAGGACACATTCCAAATGCCTTGGTATTATTAAAAGCCCT TCATGATGTGGCCCCATCTTTTATGACTTTTCCTTTTCAACTGTGCCCTCTAGCCTTATT TGATTTCTCTCAAATTCTTAAACACAGCATGCTTCACTGACCTTTAAGCCTTTGCACATA CAGTGTTGATGTGGAGCTTCCTGACCAACTCCTAATTCTCCTTCAGGCCTCAATTTAAAC ATCACTTCCTCTGGGAAGCTTTCTATTATTCCCAAGGTACTGGGATATGTTCTTGCACAG CATGCTGGGCTAATGTCACAATGGCTACCTTGTTTTATTGTTAGTATTTGATCAGCGACA CCTTGCCAGGGAGCCCCTGAGTATTGTCTGAGCAGAAACTATGGCTATCTTGTCCCCTGT AAAAAGTGTTACATTAATAAACACACACACATACATACAAAGAAATACCTGTCTTTCTCC ATATCTCAAGATCATGCTGAAAAGCCAGCATTCATGAACAAATTCCTGTGCGAAGATTGA GAATGAAAGATGAATAAGAGGTATCTTTAGAACCCAATTATGGCTGCCGTTGTTCCCTGA GTGTGAGGCTTGCTGTTAGAGTGACAGAAGGAATTTTGACTACTCAAGACCATACAAATT TGGAAATGACTCCAAAGTAAACATGGTTAGATAACTACACATTCCATTCCCCCTTTTTTA TTTCTATAGAATCCCAACTTTGTTCAAGTAGTAACATGCCCAGCTTCAGAAATGAGTCAT GATTTTTCTAAAGCAACAATATCAATCTTCTTTCCCTTCCCCAGTGATTGGTATGGAAGT TTTTGCTTTCTGCTGTAAATCAAAAGCAGAAACAGGAGAAGATTCTTTTGGGCCTCTTTC CCTCTTCCTGGCGTGGAAGTAGTTGTGAGAGCATATGATACCCAAAGTTTCGGTAGACAT TTTATAATTATGTGATGAATAACCTAAGGATAATTAAACATATAAAAGAATGGAGAAAGA GATTAAGTGATGTTATCTTCCTTTAAGGCAATCAAAATGCATCTGACAAATGGCCATCTA ATTTAAAATTCCAACTATGTAGACATCTCAAAACAAAGTCAGTATCTCAAAAAATATACTA CAAAAATTCTCATGTGTCCATTGGGGATAACTTCCAATGCTCTTTCATTGGTATTGTAGC TATGGCATTTGATTTCCAATTGTATGTGGATCAGGTAGTTGCAGGGTGACTCTCAAGGGC AACAAGACAAAGTCAAACCCTAGGTAGAAATAAGAAGGAGCTAGTACAGAAAGCAAATGC GCAGCACATAGAATCTTGGGGTTTCAGGGATATTGTTTATGAAAGGTTAGAATAGGCAAC AATTACCTGTCATATGGGTCTTGTCATTTATTATAATTTAAGGAGAATTAAAACTGAAC TAGTTGCTGGGGAGTGACATCAGCAAGATGGAGATATAGAAATCTTCAGGACCTCCTTCC GTCCATGGAACCACTGACTCAAAAATGACAAATGGAAAAAATTTACTTTCTGAGAAATCA AGAAGCCAGTTAAGAGGCTCCTGTATCTCAGATGAGTGCAAAGCCAGCTGCAACAGAGCC AGCAGAAAATTTGTTGTACTCACTCTTCATGGTCACTTCTGGCATAGCACAGTGCAATCT AGAAGAAATTCTCGGCTCCTGACTACTTTCTTGGAAAAGAAAAGAGAAAAATGTACCATAT GTCTAATATTCTGATGGGGATGGGGTGTGGGCTCCAAAGGACTAGCTTCCGTCATGCC TAAATACAAGTGCTAATTGGGAAGTCCACAATGTTGGGGGCTGCAGAAAACAAGGGCCAAC AGTTTGGACTAGCATGCACTCATTTGCCGCAGTTCCTCCTCTCACTTCATAGAATGAGTA GAAGAACCCTTAACTCTCAAGGTTTTTTTCCTGGGGAGAAAAGAGTCAAAGCAATTATA

FIG. 1S

20/51

ACCAATCTCAGAGTGCAGATGGAACCTAGCATATTCTAGATGCCTGGGGGCCATTGAGAA CAAAAGAGAGCTAGGCAACTTTCAGCAGCTCCAGAAGAACTGTGGTACCACAGATAGACA AGAATTTACACACACTGGTACAGATAAGATGAATTTGCAAAAAAAGAATAGAGGCCCCAG AATTTCTAGCTGGGTTTTTTGGTGAAGGCCTTTCTCTGTATCAAGCTAGTCCCTAAAGAC GTCACCCAGACTTGAGTGCAGTGATGCAATCATAACTCACTGCAGCCTCAAACTCCAAGG GTCAAGTGATCTTTCCACCTCAGCCTCCTGAGTAGCTGAGACTAGAGACACATGCCACTG TCAGGCTGGACTTGAACTATTGACTTCAAGGGATCCTCCTGACTCAGCCTCCCAAATCAT TGGGATTACAGGCATGACCACCATGCCTGACCTGTTTTGTTTTAAAAAAACTCAG AAAAATTTCAAAATAGCAATTATAAAGACAATGAGCTTAGAAAACCAATTAATGGACAAA ATGTAACTATAAGTAAAGAGATACATGTAAAAAGAATCAAAACAAAATTTGCAGTGGAAGA ATATGATAACCAAATTGAATATTACATTAGAGGAGTTTAATACTAGATTTGAACAAGCAG AAGAAGACTAAAAAAGAGTGAAGAAACCCTAAGGACATCATCAAGTAGACCAATATGTGT TATCAGAGTTTTAGAAGAAAAAGACAGAAAAATAGGCATAAAGCATCATTGACAAAATAA TGACCCAAAACCTCCCAATTATGAAAGACAATAGATATTCTGAATCCAGAGCACAATGGC CTGCAACTAAGATGAACCCAGAAAAGTCTATACTTCAGCACATTATAATCTAATTATCAA CAAGGGCTGTCATGAGAATATCAGCAGATTTCTCAGCAGAAAACTTGCAAAACAGAAATA AGTGGGATTACATATTCAAAGAGCTGAAAAAAAGTCTGCCAACAAAAAATCCTTTATCCA GAAGAATTTTCTTCAAAATGAAGGAGAATAAAGGATATTCCAGATAAACAAAAGCCAAGG GAATCCATCACAATTAAACCTGCCTTACAAGAAATGCTAAATGAAGTTGTTCAAGTTGAA ATAAAAGAACGCTGAACAGCAACACAAAAGCATATAAAAGTATAAAGCTCATTGGTCAAA GATAGATATAAAGGAAAAACAACGGGATATTATAATGGTGGTGGGTAACTTACTCTTCAT CCTGGTATAGAAGTTAAAAAAACCACAAGTATTAAAATAACTGTAACTATAAAATTATT **AATGAATACACAATGTAAAAATATGTAATTTGTGATACCTGATAACATACCATGTGTGGAG** GGGAGAAGTCAAAGTGTAGAGTTTTAAATAAGACTGAGGTTAGGTTTTTATCACCTTAAA ATAGATTGTTATAATATGTTTGATTTAAGCCCCATGGCAACTACAAAGAAAATACCTACA GGTAATAAACAAAAGAAATGAGAAAGAAATGAAAGTGTGTCTCAGTCCATTTTTATTTT $\tt TTCTGGAGGCTGTGAAGTTCAAGACTGAGTTGCTGCCTCTGTTGAGGGGCCTTCTTATTG$ CATCATAACATGGCAGAAGGCATCACATGACAAAAAAGCAACAGCAAGAGCCAAACTGGC TTTTATCATAGGCCTAGTTTGTGACACCTTACATAGTCCTATGAAAACCCATTAAGCCAT TAGCCCATTAATCCATTAATTCATGAATAGATTAATACATCCATGTGGGGAAAGCCCTCA TGACTCAAACCTTTCTCAAAAAACCCATCTCTTAATACTGTTACATTAGTATTAAGTTTT AACATGAGTTTCAGAGTCTAGAAATATTCACACCATAGCCTTTCACCCATGACCTCCCAT AATTTATGTCCTTATCATATGCAAATACCTTCATTCCATTCCCGTAGCCCCGAAGTCTTA ACCTGTTCTAGCACCAACTCTAAAATACGAAGTCAAGAGTCTCATCTGAGACTCAAGGCA TGATCCATCCTTGGGCAGGTTCCCTTTCAGTTGTGAAATCAAAACAAGTCATATAATTCT AAAATACAGTGCTGGTACAGGAATAAGACAGACATTCCCTTGTCGAAAGGGAAAATAAAC TAGAAGAAGGGGTTAATGGTCCCCAAGCAAGTCTTTAACACAGCAGGGCACATATTAAAT TGTAAAGCTAAAGAATACTCTTTTTTGGGTCCATGTTAAGCATTCTCTGCACAATGTGGG GAACACATTGAGCCACTCTGCCCCTATGGCTTTGCTGTGCTCAGAACACACTTCAGCTTT CTCAGATTGGAATTGCTCATTGGTGCCTGCAGCTTTCCCAGGTGGGCACTGCACACTGCT GGTGTTTCTATAATTCTAGGATCTCAAAGGCAGCTCTGGCTCTCACCCCGTATTTTTACT CAACATTGCTGTAGTGGGGCTCTCAGCCATGGCTCTGTCCCTGTGACAAGTCTCTGCCTG GGTCCCCATGCTTTTAGATACATCCTCTGAAGTCTAGGTGAAGGCCATAGTGGCCCTACA ACTCTTGCATTCTGTATCCCTGCAGAATTAGCACCAGGTGGACACTGCCAAGGCTTATGG $\verb|CTTTTGCTTTCTGGAGCAGTGAGGTAAGCTACACTTGGAGCCTCTTGAGCCAGTTGGAGT|\\$ GGCTGAGGAATGATGCGCTCACATGAAGGGAGCAGAGGAGTCCTGAGCAGCCCTGGGCAG CAAGCTGTGGAGAGTACCCTGGGCCTGTCCCCTGAAACTATTCTACCCTCCTTGGCCCCT GGGCTTTTCATGAGAGGGGGCAGTCTTAAAAATATGCAAAATACTTTTCAAACATTCTCC TCATTGTCTTAATGAATAACATCTGACTCCCTTCTATCAGTGCTAATCTCTTTAGCAAGC AGTTTTGCTGTTTACATGGCTAAGCAAGCTGCAAACTTTTCAAATCATTTTGCTGTGATT CCCTTTAATTATACATCTGTCTTTAAGTCATGTTTTTTGCTCCTGAATTGGCCAAAAGTAA

FIG. 1T

CCACACAGCCAAAAGTAGCCAAACAGCATCATGAATGCTTTGCTCCTTAAAAATTTCTTC TATAAGATATTTTACTTTATTGTCAAGTCTGGCCTTCTACACAGCCCTAGAGTATGG ACACAGTTCCAGTAAGCTTTTTGCTACTTTATACCAAGTATGACCTTTATTCCAGGTTCT GATACCTTGTTCCCCCTTTCTGTCTGAAACCTCATAACGGCCTTCATTGTCTATATGTTT ACTAGTATTTTGGCCATAATCACTTAAATAATTTATAAAATGATTCAGACTTTCCCTAGT CTTCTCATCCTCTGATCCTTCACCAGAAGCACCCTTAACACTCTATTTACAGCAATATAA GATTTTTTTTGCCTGCTCCTCCAAACCCTTCCAGCCTTTGTCCATTACCCATTTCCAAAG CCACTTGCACATTTTTAGGTTGAGCATCAGCCTCACTTCTTGTTACCAAAGCCTGTATTA GGGTTCTCCAGAGAGACAAAACCAATGGGATATACAGAAGGGGATTTGTTAGGGAAATTG GCTCACACAGTTATGGAGACTGAAAAGACCAAGGTCAAGGGGACGTATCTGGTGAGAACC TTCTCATTGTATCATAACATGGCAGATGGCATCACATGCTAAAAGAGCAAGAACAATAGC CAAACTGGATTTTATAACAGACCCACTCTTGACGACTATCCTATTCCTGTGATAAGCCAT TAATCTGTGAATCCATGAGTAAATTAATCTATTCATGAGGGCTCTGCCTCTATTGTCCCT TAAAGGCCCCACTTCTTAATACTGTTACATTGGGGATGAAGTTTCAATATGGGTTTCAGA GGAGACAAACATTCAAACCATAGTGATGTCACTACAAAAAAATTAATGAAACACAAAGGA GTACAGTAAGAGAGCAAAATACAGATAAAAGTGCTATATGATATAGAAAACAATAAAA TGGCAATAGTAGGAGTTTATCTGTCAGTAGTTACTTTAGCCATAAATGAACTAAACTCAA ACAAAAGACAAAGATTAGCTGACTGGATTTAAAAAATACTATATGCTGTCTACAAGAAGT ACAAGGAGCCCACTCCAAATTTGTAGACACACATAGGATAAAATTAAAAGGATGGAAGAA AGTATTCCATGTGAATGGTAACCAGATGAGAGCAGGGCTCATTATACTTATATCGGACAA ATAAATTGTAAGTCAATAATTGTCACAAGGAACAAAGAAGGACAATATGTAATATTAAAA GAGTCAATTCACCAGAAAGATATAACAATTTTAAACATATATGTATTCAATCTTAGGGCT TTAAAATATATAAACAAATATTAATGGAACTGAAGGGAGAAAGACAGCAATACAACAATA GTAGGAGATTTTAATTCTCAGCTTTCTTTTTCTAGAGACAGAGTCTCACTCTGTCACTCA GGCTGGAGGGCAATGGTACAATCTCAGCTCACTGCAATCTCCCAGACTCAAGTG ATTCTCCCACTTCAGCCTGCTGAGTAGCTGGGACTGCAGACATGCAACACCATACCCAGC TAATTTTTTAACTTTTTGTACAGATGAAGTCTCGTATATTGCCCAGCTGGTCTTAAACTC TTGGGCTCAAGTGATCCTTCACCTGGGCCTCCCAAAGTGCTGGGATTATAGGCATGAGCC ATGCTGTGTCACAAAACATGTTTTAACAAATTTAAAAAATACAGAAATCATATCAAATATC TTTTCTGAACACAGTGGAATGAAACTATAAATCAATTATAAAAGGAAACTGGCAATTTCA CCAATATGTGTACATTAAACAATAAATTCTTGAACAGTCCATGAGTCAAAGAAGAAATTA TAAGGGATATTTGAAATGTTTCAAGATAAATGAAAATGTCTCAAGATGAAATAAAAAGAC AACATATCCAAATTTATGGAATGCAACAAAAGTGGCAAGAGTTAAGTTTATAGTGGTAAG TGACTACATTATAAAAGAAAAAGATTTTAAGTAAACAACCTAACTTTACACCTCAGAAG CTTTTTAAAGATCAATAAAATTTACAAACCTTTGGCTAGAATAACTAAGAAAAAAGAGAG AAGACTCATAAATAATATTGTAAATAAAAAAGGAGCTATTGCAATCAAAGAGGCAGGAAC AATAAAGATTTTCAGGCTATTCTGTATAATTATACACTAACAAATTGGATAACCTAGAAG AAATGTATAAATTCTCAGAAATACACAACCTACCAAGACTGAATCAAGAAGAAATACAGA ATCTGAACAGATCTGTAACTAGTAAGGAGATTAAATCAATGATCAGAAACTTCCCAAAAA AGAAAATCCCAGGATCAGAAAACTTCACTGGAGAATTCTGCCAACATTTAATAGAAAAAA AAATGCCAATTCTTCTCAAACTTTTGCAAAAAATTGAAGAGGACGAAGCATTTCAAACTC ATTTTATGAGTCCAGCATTTTCCTGATACCAAAATGAGATAAAGATATTACAACGAACAC ACACACTTTCAAACAAGCTACAGGCCACTATCTCTGATGAATGTAAATGCAAAAGTTGTC AATAAAAATAGCAAACTGAATTCAACAGTGCATTAAAAGGATCACACACTGTGACCAAG CTATATTAACAGAACAAGGGATAAGATCACATGATAATCTCTATAAATGCTGAACAATCA TTTGACAAAGTTTAATACCCTTTCGTAATAAAAATACTCAACAAACTATGAATAGAAGGC ATGTACCTCAACACAATAATAAAGGTCACATATCAAAAGCTAACAGATAACATCATACTC AATGGTAAAAACTGAAAGCTTTTCCTCCAAGATCAGGAACTAGGTAAGAATGTCCATTCT TGCCATTTCTCATCAACGTATTACTAGAAGTCTTTGCTAGAACAATTATGCAAGAATAAG AAATAAAAAGCACTGAAATCAGCAAGGAAGAGGGAAAATTATCTCTATTCCCAGATATAA TAATCTTATATGTAGAAAATTCTAAAAATCACACAAGGAAACTGTTGCAACTAGTAAGTT

FIG. 1U

CATCAAAATTGCAGAACATAAAATCGAAATGCAAAAATCAGTTATGTTTCTATACAATAG CAGCAAACTCTCTGAAAAAGACATTACAATCCCACTTACAATATTATCAAAAATGACTAA AATGTTTAGTAATAAGCTTAACCAAGGAGGCTAACGACTTATACACTGAAAACCATAAAA GCATTACCAAAAAATAATTTTAAAAGACACAAATAAATAGAAAGATAATTCTGTTTTCAT GGGTTAGAAAACTCGATATTGTTAAAATGTGCACACTGCTGAAAGCAATTTATAGATCCT CCATGGATACTTAGAAAATCTGGAGAAAGAAGAGGCAAAGTAGAGGGTCTCATGCTTCCTG ACTTCAAAACATATTCCAAAGCCATTGTAATAGAAACAGTTTAGCACTGGCATAAAGACA **AAATAATATACAAAGCACAAAGACTATGGACAGGATAGTCTCTTCAACAATTGTGTTTGGG** AAAACTAGATAGCCATATTCAAAGGACTGAAATTAGACCCTACTCAAAAAATCAAGTCAA AATGAATTAAAAATTAAAGATCTGGGCCGGGCGTGGTGGCTCACGCCTGTAATCCCAGCA $\tt CTTTGGGAGGCCAAGGGGTCAGATCACGAGGTCAGGAGATCGAGACCATCCTGGCTAAC$ ACAGTGAAACCCCGTCTCTACTAAAAATACAAAAAATTAGCCGGGCGTGGTGGTGGGCGC CTGTAGTCCCAACTACTCAGGAGGCTGAGGCAGGAGAATGGCGTGAACCTCAGAGGCAGA GCTTGCAGTGAGGTGAGATCACGCCACTGCACTCCAGCCTGGGGGACAGAGCAAGACTCC GAGAAAAGTTTTATACCATTGGTTTTGGCAATAATTTCTTGTATACGACACCAAAGAACA GGCAGTAAAAGCAACAAAAATAGATAAGTGGAACTACATAAAATTAAAAACTGATGCAC AGAAAATAAATAAAAGAAAAAACAGAGTGTAAAAGCAAACCATGAAATGGGAGAGAATA TTTGCAAACCATATATCTGATAATGGGTTAGTATTCAAAATATATAAGGAACACCTACAA CTCAATAGCAAAAACTAACCCAATTAAAAATGGACAATGGACCTGATGGATATCTCTCC AAAGAAGATGTAAAAACAGCCAACAGATACATGAAGAGTGCTTAACATCATTAGTAATTA GGGAAATGCAAACCACATGAGCTATCATCTTACACCTGGTAGGATGACCATTATG AAACAAAAGAAAAGAGAATTAAAAAAAAAAAAGTGTTGAAAGGGATGTGGAGAAACTAGAA CCTTTGTACAGCCACTGTGAAAAATGTTTGGAAGTTCCTCAAAAAAATTAAAAATAAAA CTATACGATCCAGTAATCCCACTTTTAGATACTTTTCCAAAATATTTGAAAACAGGAACT CAAAGAGATATTTGCACTCTCATGTTTATTGTAGCCTTATTTACAATAGTCAAGAGGTGG AAACAAATGAAATATATAATGACAGATGAGTCAATAAAATGTGGCATGTACATATCATGG AATATTATTCAGCATTACAAAAGAAGAAAATCTTATAATATGCTGCAACATAGACAAACC TTGAGGACCTTATACTAAATAAAATAAACCAGTCACAGAATGACAAATACTGCATGAATA TACTTCTATGAAGTATCTAAAGTAGTCAGTCATAGAAGCAGGAAGCAGAACGGCAGCTGC ${\tt CAGGTCCTGGGAGTAAGAGTAAGAGGGAAAGTTGCATTTCAGTGGGTATAGAGTTTAAAGC}$ ATGCAAGATGAAAAAGCTCTAAAGATCTGATGTACAATAATATGCATATAATGAACAATA TTGTACTGTTCACTTAAATATGTGTTAGGTCCATGTTATGTGATTTTTACCACATTTTTT TGAAAGCAAGTTGCTAAAGAATTTGCCAAATGGAATTATAGTGACACGAGTTCAAATAAA ATTAAAAAACGAGAAACAGTAGAGTTTACTTAATTTGTTAATATATCCATATTATCATTT TAGGGAATTTTTACTAAAGCAGAGTATATAAACTATCTTTTTTTGTTCTAATGATCCATT TGTTTTAGTTTGTTTCCCATTTTTATGTAGCTAGACTGCCAGTTAATCTCCTAAAATTAT TGGCACCATATTTCCCATTTTTTCTGGCTTTTTTATTAGTAACTGGGATCCTTGCAGCTG TATCTATGTGATGCCAAACAATTAGGTTGATCAATTCTGTGACAACAAGCCATCTGGTTA CTTTAGTGAATAGGCCCTTACTTACCTTTCATAAGTTGATTCTATTCTCCTTTGTGCCTT CTCTTTAAATTACCATTATCCTGTAACCATAAATTAAAAATACAGCATCGCTTTTAAAAAC GACCCTAATTAGCATTTAGGAACAAACTACACTTGCAAAATTATTTTCGATTGGTAGAGG GAAGAAAAGGGTCTTTTTATTACTATGTATTTGTAATTACTTTTGTCACTTATGTTATTC TTGTGTCTAAATTCAACTCTAGATTTATTCTCTGTTGATATTTTTTATCACTTGAGAATA TTTTAGTTTTTCAACCTCTATATGGCGGGCTATCACTCCAAATTTAGGTTAAACTGTAGG AGTAGTTATATTTCAGTACTTCTTTTACCTAATCAGCCATTTTAAAATAATTTTGTTCAT GGCAGCAATCTGCATGACAAATTTCTACTTAATAAGCAATGAAATAGTTGGATAAATGTG TATTTCTACATGGGTGAATTTCCCAAAATTCACACTTCAAAGACAGTTGCTGACATTTTT TCAATGAGAGATTTTATTAGATAATGAGTCATCTTAGAGTTATCTTGTAAGTATTCTTTA TGTTTTATAAACATTAGAAATTAAATAGGACTACCATATGGTCTAGCAATCACACTTCTG GGTATATATCCAAAGAAAATCAGTTCAGTATGTCAAAGAGATGTTTCGTATTCATTGCAG

FIG. 1V

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization International Bureau



(43) International Publication Date 3 May 2001 (03.05.2001)

PCT

(10) International Publication Number WO 01/30991 A3

- (51) International Patent Classification7: C07K 14/705. C12Q 1/68
- (21) International Application Number: PCT/US00/23021
- (22) International Filing Date: 22 August 2000 (22.08.2000)
- (25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

09/426,290

25 October 1999 (25.10.1999)

(63) Related by continuation (CON) or continuation-in-part (CIP) to earlier application:

US

09/426,290 (CIP)

Filed on

25 October 1999 (25.10.1999)

- (71) Applicant (for all designated States except US): DECODE GENETICS EHF. [IS/IS]: Lynghals 1, IS-110 Reykjavik (IS).
- (72) Inventors; and
- (75) Inventors/Applicants (for US only): OLAFSDOTTIR, Berglind, Ran [IS/IS]; Eskihlid 15, IS-105 Reykjavík (IS). GULCHER, Jeffrey [US/US]; Unit M, 130 South Canal Street, Chicago, IL 60606 (US).

- (74) Agents: CARROLL, Alice, O. et al.: Hamilton, Brook, Smith & Reynolds, P.C., 530 Virginia Road, P.O. Box 9133, Concord, MA 01742-9133 (US).
- (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ. BA. BB, BG, BR. BY, BZ, CA, CH, CN, CR, CU, CZ, DE. DK. DM. DZ. EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (regional): ARIPO patent (GH. GM. KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT. BE. CH. CY, DE. DK. ES, FI, FR, GB, GR, IE, IT. LU, MC, NL, PT. SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

with international search report

(88) Date of publication of the international search report: 6 December 2001

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: HUMAN NARCOLEPSY GENE

(57) Abstract: The gene for hypocretin (orexin) receptor 2 (HCRTR2), which is associated with narcolepsy, is disclosed. Also described are methods of diagnosis of narcolepsy, pharmaceutical compositions comprising nucleic acids comprising the HCRTR2 gene, as well as methods of therapy of narcolepsy.

INTERNATIONAL SEARCH REPORT

Internat. I Application No PCT/US 00/23021

TDC 7 CO7K14/705 C1201/69	A. CLASSI	SIFICATION OF SUBJECT MA	ATTER ,
1FC / CO/K14//OS C12Q1/O	IPC 7	C07K14/705	C12Q1/68

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

 $\begin{array}{ll} \mbox{Minimum documentation searched (classification system followed by classification symbols)} \\ \mbox{IPC 7} & \mbox{C07K} & \mbox{C12Q} \end{array}$

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, WPI Data, BIOSIS

C. DOCUM		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 96 34877 A (HUMAN GENOME SCIENCES INC.; LI YI (US); ROSEN CRAIG A (US); SOPPET) 7 November 1996 (1996-11-07) the whole document	1-7
Y	LIN LING ET AL: "The sleep disorder canine narcolepsy is caused by a mutation in the hypocretin (orexin) receptor 2 gene" CELL,CELL PRESS, CAMBRIDGE, NA,US, vol. 98, no. 3, 6 August 1999 (1999-08-06), pages 365-376, XP002153571 ISSN: 0092-8674 abstract; figure 6	1-7

	<u>1</u> ~
* Special categories of cited documents: *A* document defining the general state of the art which is not considered to be of particular relevance	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
 "E" earlier document but published on or after the international filling date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filling date but later than the priority date claimed 	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "&" document member of the same patent family
Date of the actual completion of the international search 22 March 2001	Date of mailing of the international search report 1 9. 04. 01
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentiaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Authorized officer Hardon, E

1

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

INTERNATIONAL SEARCH REPORT

Internat. J Application No PCT/US 00/23021

	DOOLHIERT CONCIDENT TO BE STORY	PCT/US 00/23021
.(Continua ategory °	ation) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with Indication where appropriate, of the relevant passages	Relevant to claim No.
Y	SAKURAI T ET AL: "Oxerins and oxerin receptors: A family of hypothalamic neuropeptides and G Protein-coupled receptors that regulate feeding behaviour" CELL,CELL PRESS, CAMBRIDGE, NA,US, vol. 92, 20 February 1998 (1998-02-20), pages 573-585, XP002105412 ISSN: 0092-8674 page 585, column 2; figure 2	1-7
Y	ALDRICH, MICHAEL S. ET AL: "Narcolepsy and the hypocretin receptor 2 gene" NEURON (1999), 23(4), 625-626, 1999, XP000973742 the whole document	1-7
Y	SIEGEL, JEROME M.: "Narcolepsy: A key role for hypocretins (orexins)" CELL (CAMBRIDGE, MASS.) (1999), 98(4), 409-412, 20 August 1999 (1999-08-20), XP000941943 the whole document	1-7
A	LECEA L ET AL: "The hypocretins: hypothalamus-specific peptides with neuroexcitatory activity" PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA, NATIONAL ACADEMY OF SCIENCE. WASHINGTON, US, vol. 95, January 1998 (1998-01), pages 322-327, XP002105411 ISSN: 0027-8424 the whole document	1-7
Т	PEYRON CHRISTELLE ET AL: "A mutation in a case of early onset narcolepsy and a generalized absence of hypocretin peptides in human narcoleptic brains" NATURE MEDICINE, NATURE PUBLISHING, CO, US, vol. 6, no. 9, September 2000 (2000-09), pages 991-997, XP002153570 ISSN: 1078-8956	1-7

INTERNATIONAL SEARCH REPORT

Inten ational application No. PCT/US 00/23021

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)
This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1. X Claims Nos.: 7 because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy
Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)
This International Searching Authority found multiple Inventions in this international application, as follows:
1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remark on Protest The additional search fees were accompanied by the applicant's protest. No protest accompanied the payment of additional search fees.

2122		IONAL SEARCH			PCT/US	Application No 00/23021
Patent document cited in search report	t	Publication date		Patent family member(s)		Publication date
WO 9634877	A	07-11-1996	CA	22200	36 A	07-11-1996
			AU	7152		20-01-2000
			AU	24707		21-11-1996
			EP	08287		18-03-1998
			JP	115051	10 I	18-05-1999
•						

(19) World Intellectual Property Organization International Bureau



| 1917| | 1917| | 1917| | 1917| | 1917| | 1917| | 1917| | 1917| | 1917| | 1917| | 1917| | 1917| | 1917| | 1917

(43) International Publication Date 3 May 2001 (03.05.2001)

PCT

(10) International Publication Number WO 01/30991 A2

(51) International Patent Classification7: C12N 15/00

(21) International Application Number: PCT/US00/23021

(22) International Filing Date: 22 August 2000 (22.08.2000)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

09/426,290

25 October 1999 (25.10.1999) US

(63) Related by continuation (CON) or continuation-in-part (CIP) to earlier application:

US Filed on 09/426,290 (CIP) 25 October 1999 (25.10.1999)

(71) Applicant (for all designated States except US): DECODE GENETICS EHF. [IS/IS]; Lynghals 1, IS-110 Reykjavik

(72) Inventors; and

(75) Inventors/Applicants (for US only): OLAFSDOTTIR, Berglind, Ran [IS/IS]; Eskihlid 15, IS-105 Reykjavik (IS). GULCHER, Jeffrey [US/US]; Unit M, 130 South Canal Street, Chicago, IL 60606 (US). (74) Agents: CARROLL, Alice, O. et al.; Hamilton, Brook, Smith & Reynolds, P.C., Two Militia Drive, Lexington, MA 02421 (US).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

 Without international search report and to be republished upon receipt of that report.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: HUMAN NARCOLEPSY GENE

(57) Abstract: The gene for hypocretin (orexin) receptor 2 (HCRTR2), which is associated with narcolepsy, is disclosed. Also described are methods of diagnosis of narcolepsy, pharmaceutical compositions comprising nucleic acids comprising the HCRTR2 gene, as well as methods of therapy of narcolepsy.

-1-

HUMAN NARCOLEPSY GENE

RELATED APPLICATION

This application is a Continuation-in-Part of U.S. Serial No. 09/426,290, filed October 25, 1999, the entire teachings of which are incorporated herein by reference.

BACKGROUND OF THE INVENTION

Narcolepsy, a disorder which affects approximately 1 in 2,000 individuals, is characterized by daytime sleepiness, sleep fragmentation, and symptoms of abnormal rapid eye movement (REM) sleep that include cataplexy (loss of muscle 10 tone), sleep paralysis, and hypnagogic hallucinations (Aldrich, M.S., Neurology 42:34-43 (1992); Siegel, J.M., Cell 98:409-412 (1999)). In humans, susceptibility to narcolepsy has been associated with a specific human leukocyte antigen (HLA) alleles, including DQB1*0602 (Mignot, E., Neurology 50:S16-22 (1998); Kadotani, H. et al., Genome Res. 8:427-434 (1998); Faraco, J. et al., J. Hered. 90:129-132 15 (1999)); however, attempts to verify narcolepsy as an autoimmune disorder have failed (Mignot, E. et al., Adv. Neuroimmunol. 5:23-37 (1995); Mignot, E., Curr. Opin. Pulm. Med. 2:482-487 (1996)). In a canine model of narcolepsy, the disorder is transmitted as an autosomal recessive trait, canarc-1 (Foutz, A.S. et al., Sleep 1:413-421 91979); Baker, T.L. and Dement, W.C., Brain Mechanisms of Sleep (D.J. McGinty et al., eds., New York: Raven Press, pp. 199-233 (1985)). The possibility of linkage between canarc-1 and the canine major histocompatibility complex has been excluded (Mignot, E. et al., Proc. Natl. Acad. Sci. USA 88:3475-3478 (1991)).

10

15

A mutation in the hypocretin (orexin) receptor 2 gene in canines has been identified in narcolepsy (Lin, L. et al., Cell 98:365-376 (1999));

Hypocrexins/orexins (orexin-A and -B) are neuropeptides associated with regulation of food consumption (de Lecea, L., et al., Proc. natl. Acad. Sci. USA 95:322-327 (1998); Sakurai, T. et al., Cell 92:573-585 (1998)) as well as other possible functions (Peyron, C. et al., J. Neurosci. 18:9996-10015 (1998)). Human cDNA of receptors for orexins have been cloned (Sakurai, T. et al., Cell 92:573-585 (1998)), however, full human genes for the orexin receptors have not yet been identified.

Diagnosis of narcolepsy is difficult, as it is necessary to distinguish narcolepsy from other conditions such as chronic fatigue syndrome or other sleep disorders (Ambrogetti, A. and Olson, L.C., *Med. J. Aust. 160*:426-429 (1994); Aldrich, M.S., *Neurology 50*:S2-7 (1998)). Methods of diagnosing narcolepsy based on specific criteria would facilitate identification of the disease, reduce the time and expense associated with diagnosis, and expedite commencement of treatment.

SUMMARY OF THE INVENTION

As described herein, a full gene for the human hypocretin (orexin) receptor 2 (HCRTR2) has been identified. The sequence of the HCRTR2 gene as described herein is shown in Figure 1 (SEQ ID NO: 1). Accordingly, this invention pertains to an isolated nucleic acid molecule containing the HCRTR2 gene. The invention also relates to DNA constructs comprising the nucleic acid molecules described herein operatively linked to a regulatory sequence, and to recombinant host cells, such as bacterial cells, fungal cells, plant cells, insect cells and mammalian cells, comprising the nucleic acid molecules described herein operatively linked to a regulatory sequence. The invention also pertains to methods of diagnosing narcolepsy in an individual. The methods include detecting the presence of a mutation in the HCRTR2 gene. The invention additionally pertains to pharmaceutical compositions comprising the HCRTR2 nucleic acids of the invention. The invention further pertains to methods of treating narcolepsy, by administering HCRTR2 nucleic acids

of the invention or compositions comprising the HCRTR2 nucleic acids. The methods of the invention allow the accurate diagnosis of narcolepsy and reduce the need for time-consuming and expensive sleep laboratory assessments.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1A to Fig. 1AY depict the sequence of the human orexin receptor 2 gene (SEQ ID NO:1) and the encoded receptor (SEQ ID NO:2).

The foregoing and other objects, features and advantages of the invention will be apparent from the following more particular description of preferred embodiments of the invention, as illustrated in the accompanying drawings

10 DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to a human hypocretin (orexin) receptor 2 (HCRTR2) gene, and the relationship of the gene to narcolepsy. As described herein, Applicants have isolated the HCRTR2 gene. The gene and its products are implicated in the pathogenesis of narcolepsy, as mutations in a closely related receptor, hypocretin (orexin) receptor 2, have been associated with the presence of narcolepsy in a well-established canine model of narcolepsy (Lin, L. et al., Cell 98:365-376 (1999)).

NUCLEIC ACIDS OF THE INVENTION

Accordingly, the invention pertains to an isolated nucleic acid molecule

containing the human HCRTR2 gene. The term, "HCRTR2 gene," refers to an
isolated genomic nucleic acid molecule that encodes the human hypocretin (orexin)
receptor 2. As used herein, the term, "genomic nucleic acid molecule" indicates that
the nucleic acid molecule contains introns and exons as are found in genomic DNA
(i.e., not cDNA). The nucleic acid molecules can be double-stranded or singlestranded; single stranded nucleic acid molecules can be either the coding (sense)
strand or the non-coding (antisense) strand. The nucleic acid molecule can
additionally contain a marker sequence, for example, a nucleotide sequence which
encodes a polypeptide, to assist in isolation or purification of the polypeptide. Such

5

15

30

sequences include, but are not limited to, those which encode a glutathione-S-transferase (GST) fusion protein and those which encode a hemagglutinin A (HA) peptide marker from influenza. In a preferred embodiment, the nucleic acid molecule has the sequence shown in the Figure (SEQ ID NO:1).

As used herein, an "isolated" or "substantially pure" gene or nucleic acid molecule is intended to mean a gene which is not flanked by nucleotide sequences which normally (in nature) flank the gene (as in other genomic sequences). Thus, an isolated gene can include a gene which is synthesized chemically or by recombinant means. Thus, recombinant DNA contained in a vector are included in the definition of "isolated" as used herein. Also, isolated nucleotide sequences include recombinant DNA molecules in heterologous host cells, as well as partially or substantially purified DNA molecules in solution. Such isolated nucleotide sequences are useful in the manufacture of the encoded protein, as probes for isolating homologous sequences (e.g., from other mammalian species), for gene mapping (e.g., by in situ hybridization with chromosomes), or for detecting expression of the HCRTR2 gene in tissue (e.g., human tissue), such as by Northern blot analysis.

The present invention also encompasses variations of the nucleic acid sequences of the invention. Such variations can be naturally-occurring, such as in the case of allelic variation, or non-naturally-occurring, such as those induced by various mutagens and mutagenic processes. Intended variations include, but are not limited to, addition, deletion and substitution of one or more nucleotides which can result in conservative or non-conservative amino acid changes, including additions and deletions. Preferably, the nucleotide or amino acid variations are silent or conserved; that is, they do not alter the characteristics or activity of the hypocretin (orexin) receptor 2.

Other alterations of the nucleic acid molecules of the invention can include, for example, labeling, methylation, internucleotide modifications such as uncharged linkages (e.g., methyl phosphonates, phosphotriesters, phosphoamidates, carbamates), charged linkages (e.g., phosphorothioates, phosphorodithioates), pendent moieties (e.g., polypeptides), intercalators (e.g., acridine, psoralen),

-5-

chelators, alkylators, and modified linkages (e.g., alpha anomeric nucleic acids). Also included are synthetic molecules that mimic nucleic acid molecules in the ability to bind to a designated sequences via hydrogen bonding and other chemical interactions. Such molecules include, for example, those in which peptide linkages substitute for phosphate linkages in the backbone of the molecule.

5

10

15

20

25

30

The invention also relates to fragments of the isolated nucleic acid molecules described herein. The term "fragment" is intended to encompass a portion of a nucleic acid sequence described herein which is from at least about 25 contiguous nucleotides to at least about 50 contiguous nucleotides or longer in length. One or more introns can also be present. Such fragments are useful as probes, e.g., for diagnostic methods, as described below and also as primers or probes. Particularly preferred primers and probes selectively hybridize to a nucleic acid molecule containing the HCRTR2 gene described herein.

The invention also pertains to nucleic acid molecules which hybridize under high stringency hybridization conditions, such as for selective hybridization, to a nucleotide sequence described herein (e.g., nucleic acid molecules which specifically hybridize to a nucleic acid containing the HCRTR2 gene described herein). Hybridization probes are oligonucleotides which bind in a base-specific manner to a complementary strand of nucleic acid. Suitable probes include polypeptide nucleic acids, as described in (Nielsen *et al.*, *Science* 254, 1497-1500 (1991)).

Such nucleic acid molecules can be detected and/or isolated by specific hybridization (e.g., under high stringency conditions). "Stringency conditions" for hybridization is a term of art which refers to the incubation and wash conditions, e.g., conditions of temperature and buffer concentration, which permit hybridization of a particular nucleic acid to a second nucleic acid; the first nucleic acid may be perfectly (i.e., 100%) complementary to the second, or the first and second may share some degree of complementarity which is less than perfect (e.g., 60%, 75%, 85%, 95%). For example, certain high stringency conditions can be used which distinguish perfectly complementary nucleic acids from those of less complementarity.

20

25

-6-

"High stringency conditions", "moderate stringency conditions" and "low stringency conditions" for nucleic acid hybridizations are explained on pages 2.10.1-2.10.16 and pages 6.3.1-6 in Current Protocols in Molecular Biology (Ausubel, F.M. et al., "Current Protocols in Molecular Biology", John Wiley & Sons, (1998)) the 5 teachings of which are hereby incorporated by reference. The exact conditions which determine the stringency of hybridization depend not only on ionic strength (e.g., 0.2XSSC, 0.1XSSC), temperature (e.g., room temperature, 42°C, 68°C) and the concentration of destabilizing agents such as formamide or denaturing agents such as SDS, but also on factors such as the length of the nucleic acid sequence, base 10 composition, percent mismatch between hybridizing sequences and the frequency of occurrence of subsets of that sequence within other non-identical sequences. Thus, high, moderate or low stringency conditions can be determined empirically. By varying hybridization conditions from a level of stringency at which no hybridization occurs to a level at which hybridization is first observed, conditions which will allow a given sequence to hybridize (e.g., selectively) with the most similar sequences in the sample can be determined.

Exemplary conditions are described in Krause, M.H. and S.A. Aaronson, *Methods in Enzymology, 200*:546-556 (1991). Also, in, Ausubel, *et al.*, "Current *Protocols in Molecular Biology*", John Wiley & Sons, (1998), which describes the determination of washing conditions for moderate or low stringency conditions. Washing is the step in which conditions are usually set so as to determine a minimum level of complementarity of the hybrids. Generally, starting from the lowest temperature at which only homologous hybridization occurs, each °C by which the final wash temperature is reduced (holding SSC concentration constant) allows an increase by 1% in the maximum extent of mismatching among the sequences that hybridize. Generally, doubling the concentration of SSC results in an increase in T_m of ~17°C. Using these guidelines, the washing temperature can be determined empirically for high, moderate or low stringency, depending on the level of mismatch sought.

For example, a low stringency wash can comprise washing in a solution containing 0.2XSSC/0.1% SDS for 10 min at room temperature; a moderate

10

20

25

30

-7-

stringency wash can comprise washing in a prewarmed solution (42°C) solution containing 0.2XSSC/0.1% SDS for 15 min at 42°C; and a high stringency wash can comprise washing in prewarmed (68°C) solution containing 0.1XSSC/0.1%SDS for 15 min at 68°C. Furthermore, washes can be performed repeatedly or sequentially to 5 obtain a desired result as known in the art. Equivalent conditions can be determined by varying one or more of the parameters given as an example, as known in the art, while maintaining a similar degree of identity or similarity between the target nucleic acid molecule and the primer or probe used.

Hybridizable nucleic acid molecules are useful as probes and primers, e.g., for diagnostic applications, as described below. As used herein, the term "primer" refers to a single-stranded oligonucleotide which acts as a point of initiation of template-directed DNA synthesis under appropriate conditions (e.g., in the presence of four different nucleoside triphosphates and an agent for polymerization, such as, DNA or RNA polymerase or reverse transcriptase) in an appropriate buffer and at a 15 suitable temperature. The appropriate length of a primer depends on the intended use of the primer, but typically ranges from 15 to 30 nucleotides. Short primer molecules generally require cooler temperatures to form sufficiently stable hybrid complexes with the template. A primer need not reflect the exact sequence of the template, but must be sufficiently complementary to hybridize with a template. The term "primer site" refers to the area of the target DNA to which a primer hybridizes. The term "primer pair" refers to a set of primers including a 5' (upstream) primer that hybridizes with the 5' end of the DNA sequence to be amplified and a 3' (downstream) primer that hybridizes with the complement of the 3' end of the sequence to be amplified.

The invention also pertains to nucleotide sequences which have a substantial identity with the nucleotide sequences described herein; particularly preferred are nucleotide sequences which have at least about 70%, and more preferably at least about 80% identity, and even more preferably at least about 90% identity, with nucleotide sequences described herein. Particularly preferred in this instance are nucleotide sequences encoding hypocretin (orexin) receptor 2.

To determine the percent identity of two nucleotide sequences, the sequences are aligned for optimal comparison purposes (e.g., gaps can be introduced in the sequence of a first nucleotide sequence). The nucleotides at corresponding nucleotide positions are then compared. When a position in the first sequence is occupied by the same nucleotide as the corresponding position in the second sequence, then the molecules are identical at that position. The percent identity between the two sequences is a function of the number of identical positions shared by the sequences (i.e., % identity = # of identical positions/total # of positions x 100).

The determination of percent identity between two sequences can be 10 accomplished using a mathematical algorithm. A preferred, non-limiting example of a mathematical algorithm utilized for the comparison of two sequences is the algorithm of Karlin et al. (Proc. Natl. Acad. Sci. USA, 90:5873-5877 (1993)). Such an algorithm is incorporated into the NBLAST program which can be used to identify sequences having the desired identity to nucleotide sequences of the 15 invention. To obtain gapped alignments for comparison purposes, Gapped BLAST can be utilized as described in Altschul et al. (Nucleic Acids Res, 25:3389-3402 (1997)). When utilizing BLAST and Gapped BLAST programs, the default parameters of the respective programs (e.g., NBLAST) can be used. See 20 http://www.ncbi.nlm.nih.gov. In one embodiment, parameters for sequence comparison can be set at W=12. Parameters can also be varied (e.g., W=5 or W=20). The value "W" determines how many continuous nucleotides must be identical for the program to identify two sequences as containing regions of identity.

The invention also provides expression vectors containing a nucleic acid

comprising the HCRTR2 gene, operatively linked to at least one regulatory
sequence. Many such vectors are commercially available, and other suitable vectors
can be readily prepared by the skilled artisan. "Operatively linked" is intended to
mean that the nucleic acid sequence is linked to a regulatory sequence in a manner
which allows expression of the nucleic acid sequence. Regulatory sequences are artrecognized and are selected to produce a hypocretin (orexin) receptor 2.
Accordingly, the term "regulatory sequence" includes promoters, enhancers, and

other expression control elements such as those described in Goeddel, Gene Expression Technology: Methods in Enzymology 185, Academic Press, San Diego, CA (1990). For example, the native regulatory sequences or regulatory sequences native to the transformed host cell can be employed. It should be understood that the 5 design of the expression vector may depend on such factors as the choice of the host cell to be transformed and/or the receptor desired to be expressed. For instance, the gene of the present invention can be expressed by ligating the gene into a vector suitable for expression in either prokaryotic cells, eukaryotic cells or both (see, for example, Broach, et al., Experimental Manipulation of Gene Expression, ed. M. Inouye (Academic Press, 1983) p. 83; Molecular Cloning: A Laboratory Manual, 2nd Ed., ed. Sambrook et al. (Cold Spring Harbor Laboratory Press, 1989) Chapters 16 and 17). Typically, expression constructs will contain one or more selectable markers, including, but not limited to, the gene that encodes dihydrofolate reductase and the genes that confer resistance to neomycin, tetracycline, ampicillin, chloramphenicol, kanamycin and streptomycin resistance. Vectors can also include, for example, an autonomously replicating sequence (ARS), expression control sequences, ribosome-binding sites, RNA splice sites, polyadenylation sites, transcriptional terminator sequences, secretion signals and mRNA stabilizing sequences.

Prokaryotic and eukaryotic host cells transformed by the described vectors are also provided by this invention. For instance, cells which can be transformed with the vectors of the present invention include, but are not limited to, bacterial cells such as E. coli (e.g., E. coli K12 strains), Streptomyces, Pseudomonas, Serratia marcescens and Salmonella typhimurium, insect cells (baculovirus), including

Drosophila, fungal cells, such as yeast cells, plant cells and mammalian cells, such as thymocytes, Chinese hamster ovary cells (CHO), and COS cells. The host cells can be transformed by the described vectors by various methods (e.g., electroporation, transfection using calcium chloride, rubidium chloride, calcium phosphate, DEAE-dextran, or other substances; microprojectile bombardment;

lipofection, infection where the vector is an infectious agent such as a retroviral genome, and other methods), depending on the type of cellular host.

-10-

The nucleic acid molecules of the present invention can be produced, for example, by replication in a suitable host cell, as described above. Alternatively, the nucleic acid molecules can also be produced by chemical synthesis.

PCT/US00/23021

The nucleotide sequences of the nucleic acid molecules described herein

(e.g., a nucleic acid molecule comprising SEQ ID NO:1) can be amplified by methods known in the art. For example, this can be accomplished by e.g., PCR. See generally PCR Technology: Principles and Applications for DNA Amplification (ed. H.A. Erlich, Freeman Press, NY, NY, 1992); PCR Protocols: A Guide to Methods and Applications (eds. Innis, et al., Academic Press, San Diego, CA, 1990); Mattila et al., Nucleic Acids Res. 19, 4967 (1991); Eckert et al., PCR Methods and Applications 1, 17 (1991); PCR (eds. McPherson et al., IRL Press, Oxford); and U.S. Patent 4,683,202.

Other suitable amplification methods include the ligase chain reaction (LCR) (see Wu and Wallace, *Genomics* 4, 560 (1989), Landegren et al., Science 241, 1077 (1988), transcription amplification (Kwoh et al., Proc. Natl. Acad. Sci. USA 86, 1173 (1989)), and self-sustained sequence replication (Guatelli et al., Proc. Nat. Acad. Sci. USA, 87, 1874 (1990)) and nucleic acid based sequence amplification (NASBA). The latter two amplification methods involve isothermal reactions based on isothermal transcription, which produce both single stranded RNA (ssRNA) and double stranded DNA (dsDNA) as the amplification products in a ratio of about 30 or 100 to 1, respectively.

The amplified DNA can be radiolabeled and used as a probe for screening a library or other suitable vector to identify homologous nucleotide sequences. Corresponding clones can be isolated, DNA can be obtained following *in vivo*25 excision, and the cloned insert can be sequenced in either or both orientations by art recognized methods, to identify the correct reading frame encoding a protein of the appropriate molecular weight. For example, the direct analysis of the nucleotide sequence of homologous nucleic acid molecules of the present invention can be accomplished using either the dideoxy chain termination method or the Maxam
Gilbert method (see Sambrook *et al.*, *Molecular Cloning, A Laboratory Manual* (2nd Ed., CSHP, New York 1989); Zyskind *et al.*, *Recombinant DNA Laboratory*

Manual, (Acad. Press, 1988)). Using these or similar methods, the protein(s) and the DNA encoding the protein can be isolated, sequenced and further characterized.

METHODS OF DIAGNOSIS

The nucleic acids and the proteins described above can be used to detect, in an individual, a mutation in the HCRTR2 gene that is associated with narcolepsy. In 5 one embodiment of the invention, diagnosis of narcolepsy is made by detecting a mutation in the HCRTR2 gene. The mutation can be the insertion or deletion of a single nucleotide, or of more than one nucleotide, resulting in a frame shift mutation; the change of at least one nucleotide, resulting in a change in the encoded amino acid; the change of at least one nucleotide, resulting in the generation of a premature 10 stop codon; the deletion of several nucleotides, resulting in a deletion of one or more amino acids encoded by the nucleotides; the insertion of one or several nucleotides, such as by unequal recombination or gene conversion, resulting in an interruption of the coding sequence of the gene; duplication of all or a part of the gene; transposition of all or a part of the gene; or rearrangement of all or a part of the gene. 15 More than one such mutation may be present in a single gene. Such sequence changes cause a mutation in the receptor encoded by the HCRTR2 gene. For example, if the mutation is a frame shift mutation, the frame shift can result in a change in the encoded amino acids, and/or can result in the generation of a premature stop codon, causing generation of a truncated receptor. Alternatively, a 20 mutation associated with narcolepsy can be a synonymous mutation in one or more nucleotides (i.e., a mutation that does not result in a change in the receptor encoded by the HCRTR2 gene, such as a mutation in an intron or an untranslated portion of the gene). Such a polymorphism may alter splicing sites, affect the stability or transport of mRNA, or otherwise affect the transcription or translation of the gene. 25 A HCRTR2 gene that has any of the mutations described above is referred to herein as a "mutant gene." It is likely that a mutation in the HCRTR2 gene is associated with narcolepsy in humans because of the association between a mutation in the HCRTR2 gene and narcolepsy in dogs (Lin, L. et al., Cell 98:365-376 (1999), the 30 entire teachings of which are incorporated herein by reference). In a preferred

embodiment, the mutation in the HCRTR2 gene is to a deletion mutation, for example, a deletion that corresponds to the deletions found in the hypocretin (orexin) receptor 2 in narcoleptic dogs as described by Lin et al., supra (e.g., a deletion of one or more exons, such as a deletion of the fourth exon, that can be caused by insertion of one or more nucleotides upstream of the splice site of the exon, or a deletion of exon 6, that can be caused by a G to A transition in the splice junction consensus sequence). In another preferred embodiment, the mutation in the HCRTR2 gene is mutation that effects a "knockout" of the entire gene, such as deletion of the first exon as described by Chemelli, R.M. et al, (Cell 98:437-451 (1999), the entire teachings of which are incorporated herein). In a third preferred embodiment, the mutation in the HCRTR2 gene is a mutation in an intron, that affects splicing (joining of exons) during translation of the HCRTR2 gene.

-12-

PCT/US00/23021

In a first method of diagnosing narcolepsy, hybridization methods, such as Southern analysis, are used (see Current Protocols in Molecular Biology, Ausubel, F. et al., eds., John Wiley & Sons, including all supplements through 1999). For example, a test sample of genomic DNA, RNA, or cDNA, is obtained from an individual suspected of having (or carrying a defect for) narcolepsy (the "test individual"). The individual can be an adult, child, or fetus. The test sample can be from any source which contains genomic DNA, such as a blood sample, sample of amniotic fluid, sample of cerebrospinal fluid, or tissue sample from skin, muscle, 20 placenta, gastrointestinal tract or other organs. A test sample of DNA from fetal cells or tissue can be obtained by appropriate methods, such as by amniocentesis or chorionic villus sampling. The DNA, RNA, or cDNA sample is then examined to determine whether a mutation in the HCRTR2 gene is present. The presence of the mutation can be indicated by hybridization of the gene in the test sample to a nucleic acid probe. A "nucleic acid probe", as used herein, can be a DNA probe or an RNA probe; the nucleic acid probe contains at least one mutation in the HCRTR2 gene. The probe can be one of the nucleic acid molecules described above (e.g., the gene, a vector comprising the gene, etc.)

To diagnose narcolepsy by hybridization, a hybridization sample is formed by contacting the test sample containing a HCRTR2 gene, with at least one nucleic

10

15

acid probe. The hybridization sample is maintained under conditions which are sufficient to allow specific hybridization of the nucleic acid probe to the HCRTR2 gene. "Specific hybridization", as used herein, indicates exact hybridization (e.g., with no mismatches). Specific hybridization can be performed under high stringency conditions or moderate stringency conditions, for example, as described above. In a particularly preferred embodiment, the hybridization conditions for specific hybridization are high stringency.

Specific hybridization, if present, is then detected using standard methods. If specific hybridization occurs between the nucleic acid probe and the HCRTR2 gene in the test sample, then the HCRTR2 gene has the mutation that is present in the nucleic acid probe. More than one nucleic acid probe can also be used concurrently in this method. Specific hybridization of any one of the nucleic acid probes is indicative of a mutation in the HCRTR2 gene, and is therefore diagnostic for narcolepsy.

In another hybridization method, Northern analysis (see Current Protocols in Molecular Biology, Ausubel, F. et al., eds., John Wiley & Sons, supra) is used to identify the presence of a mutation associated with narcolepsy. For Northern analysis, a test sample of RNA is obtained from the individual by appropriate means. Specific hybridization of a nucleic acid probe, as described above, to RNA from the individual is indicative of a mutation in the HCRTR2 gene, and is therefore diagnostic for narcolepsy

For representative examples of use of nucleic acid probes, see, for example, U.S. Patents No. 5,288,611 and 4,851,330. Alternatively, a peptide nucleic acid (PNA) probe can be used instead of a nucleic acid probe in the hybridization

25 methods described above. PNA is a DNA mimic having a peptide-like, inorganic backbone, such as N-(2-aminoethyl)glycine units, with an organic base (A, G, C, T or U) attached to the glycine nitrogen via a methylene carbonyl linker (see, for example, Nielsen, P.E. et al., Bioconjugate Chemistry, 1994, 5, American Chemical Society, p. 1 (1994). The PNA probe can be designed to specifically hybridize to a gene having a polymorphism associated with autoimmune disease. Hybridization of the PNA probe to the HCRTR2 gene is diagnostic for narcolepsy..

-14-

In another method of the invention, mutation analysis by restriction digestion can be used to detect mutant genes, or genes containing polymorphisms, if the mutation or polymorphism in the gene results in the creation or elimination of a restriction site. A test sample containing genomic DNA is obtained from the individual. Polymerase chain reaction (PCR) can be used to amplify the HCRTR2 gene (and, if necessary, the flanking sequences) in the test sample of genomic DNA from the test individual. RFLP analysis is conducted as described (see Current Protocols in Molecular Biology, supra). The digestion pattern of the relevant DNA fragment indicates the presence or absence of the mutation in the HCRTR2 gene, and therefore indicates the presence or absence of narcolepsy.

5

10

Sequence analysis can also be used to detect specific mutations in the HCRTR2 gene. A test sample of DNA is obtained from the test individual. PCR can be used to amplify the gene, and/or its flanking sequences. The sequence of the HCRTR2 gene, or a fragment of the gene is determined, using standard methods. The sequence of the gene (or gene fragment) is compared with the nucleic acid sequence of the gene, as described above. The presence of a mutation in the HCRTR2 gene indicates that the individual has narcolepsy.

Allele-specific oligonucleotides can also be used to detect the presence of a mutation in the HCRTR2 gene, through the use of dot-blot hybridization of amplified proteins with allele-specific oligonucleotide (ASO) probes (see, for 20 example, Saiki, R. et al., (1986), Nature (London) 324:163-166). An "allele-specific oligonucleotide" (also referred to herein as an "allele-specific oligonucleotide probe") is an oligonucleotide of approximately 10-50 base pairs, preferably approximately 15-30 base pairs, that specifically hybridizes to the HCRTR2 gene, and that contains a mutation associated with narcolepsy. An allele-specific 25 oligonucleotide probe that is specific for particular mutation in the HCRTR2 gene can be prepared, using standard methods (see Current Protocols in Molecular Biology, supra). To identify mutations in the gene that are associated with narcolepsy, a test sample of DNA is obtained from the individual. PCR can be used to amplify all or a fragment of the HCRTR2 gene, and its flanking sequences. The 30 DNA containing the amplified HCRTR2 gene (or fragment of the gene) is dotblotted, using standard methods (see Current Protocols in Molecular Biology, supra), and the blot is contacted with the oligonucleotide probe. The presence of specific hybridization of the probe to the amplified HCRTR2 gene is then detected. Specific hybridization of an allele-specific oligonucleotide probe to DNA from the individual is indicative of a mutation in the HCRTR2 gene, and is therefore indicative of narcolepsy.

Other methods of nucleic acid analysis can be used to detect mutations in the HCRTR2 gene, for the diagnosis of narcolepsy. Representative methods include direct manual sequencing; automated fluorescent sequencing; single-stranded conformation polymorphism assays (SSCA); clamped denaturing gel electrophoresis (CDGE) heteoduplex analysis; chemical mismatch cleavage (CMC); RNase protection assays; use of proteins which recognize nucleotide mismatches, such as *E. coli* mutS protein; allele-specific PCR, and other methods.

PHARMACEUTICAL COMPOSITIONS

The present invention also pertains to pharmaceutical compositions comprising nucleic acids described herein, particularly nucleic acids containing the HCRTR2 gene described herein. For instance, a nucleotide or nucleic acid construct (vector) comprising a nucleotide of the present invention can be formulated with a physiologically acceptable carrier or excipient to prepare a pharmaceutical composition. The carrier and composition can be sterile. The formulation should suit the mode of administration.

Suitable pharmaceutically acceptable carriers include but are not limited to water, salt solutions (e.g., NaCl), saline, buffered saline, alcohols, glycerol, ethanol, gum arabic, vegetable oils, benzyl alcohols, polyethylene glycols, gelatin, carbohydrates such as lactose, amylose or starch, dextrose, magnesium stearate, talc, silicic acid, viscous paraffin, perfume oil, fatty acid esters, hydroxymethylcellulose, polyvinyl pyrolidone, etc., as well as combinations thereof. The pharmaceutical preparations can, if desired, be mixed with auxiliary agents, e.g., lubricants, preservatives, stabilizers, wetting agents, emulsifiers, salts for influencing osmotic

-16-

pressure, buffers, coloring, flavoring and/or aromatic substances and the like which do not deleteriously react with the active compounds.

The composition, if desired, can also contain minor amounts of wetting or emulsifying agents, or pH buffering agents. The composition can be a liquid solution, suspension, emulsion, tablet, pill, capsule, sustained release formulation, or powder. The composition can be formulated as a suppository, with traditional binders and carriers such as triglycerides. Oral formulation can include standard carriers such as pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, polyvinyl pyrollidone, sodium saccharine, cellulose, magnesium carbonate, etc.

10

Methods of introduction of these compositions include, but are not limited to, intradermal, intramuscular, intraperitoneal, intraocular, intravenous, subcutaneous, oral and intranasal. Other suitable methods of introduction can also include gene therapy (as described below), rechargeable or biodegradable devices, particle acceleration devises ("gene guns") and slow release polymeric devices. The pharmaceutical compositions of this invention can also be administered as part of a combinatorial therapy with other agents.

The composition can be formulated in accordance with the routine procedures as a pharmaceutical composition adapted for administration to human beings. For example, compositions for intravenous administration typically are solutions in sterile isotonic aqueous buffer. Where necessary, the composition may also include a solubilizing agent and a local anesthetic to ease pain at the site of the injection. Generally, the ingredients are supplied either separately or mixed together in unit dosage form, for example, as a dry lyophilized powder or water free concentrate in a hermetically sealed container such as an ampoule or sachette indicating the quantity of active agent. Where the composition is to be administered by infusion, it can be dispensed with an infusion bottle containing sterile pharmaceutical grade water, saline or dextrose/water. Where the composition is administered by injection, an ampoule of sterile water for injection or saline can be provided so that the ingredients may be mixed prior to administration.

-17-

For topical application, nonsprayable forms, viscous to semi-solid or solid forms comprising a carrier compatible with topical application and having a dynamic viscosity preferably greater than water, can be employed. Suitable formulations include but are not limited to solutions, suspensions, emulsions, creams, ointments, powders, enemas, lotions, sols, liniments, salves, aerosols, etc., which are, if desired, sterilized or mixed with auxiliary agents, e.g., preservatives, stabilizers, wetting agents, buffers or salts for influencing osmotic pressure, etc. The agent may be incorporated into a cosmetic formulation. For topical application, also suitable are sprayable aerosol preparations wherein the active ingredient, preferably in combination with a solid or liquid inert carrier material, is packaged in a squeeze bottle or in admixture with a pressurized volatile, normally gaseous propellant, e.g., pressurized air.

Agents described herein can be formulated as neutral or salt forms. Pharmaceutically acceptable salts include those formed with free amino groups such as those derived from hydrochloric, phosphoric, acetic, oxalic, tartaric acids, etc., and those formed with free carboxyl groups such as those derived from sodium, potassium, ammonium, calcium, ferric hydroxides, isopropylamine, triethylamine, 2-ethylamino ethanol, histidine, procaine, etc.

The agents are administered in a therapeutically effective amount. The

amount of agents which will be therapeutically effective in the treatment of
narcolepsy can be determined by standard clinical techniques. In addition, in vitro
or in vivo assays may optionally be employed to help identify optimal dosage ranges.

The precise dose to be employed in the formulation will also depend on the route of
administration, and the seriousness of the disease or disorder, and should be decided
according to the judgment of a practitioner and each patient's circumstances.

Effective doses may be extrapolated from dose-response curves derived from in vitro
or animal model test systems.

The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Optionally associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture.

10

use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use of sale for human administration. The pack or kit can be labeled with information regarding mode of administration, sequence of drug administration (e.g., separately, sequentially or concurrently), or the like. The pack 5 or kit may also include means for reminding the patient to take the therapy. The pack or kit can be a single unit dosage of the combination therapy or it can be a plurality of unit dosages. In particular, the agents can be separated, mixed together in any combination, present in a single vial or tablet. Agents assembled in a blister pack or other dispensing means is preferred. For the purpose of this invention, unit dosage is intended to mean a dosage that is dependent on the individual pharmacodynamics of each agent and administered in FDA approved dosages in standard time courses.

METHODS OF THERAPY

The present invention also pertains to methods of therapy for narcolepsy, utilizing the pharmaceutical compositions comprising nucleic acids, as described 15 herein. The therapy is designed to replace/supplement activity of the hypocretin(orexin) receptor 2 in an individual, such as by administering a nucleic acid comprising the HCRTR2 gene or a derivative or active fragment thereof. In one embodiment of the invention, a nucleic acid of the invention is used in the treatment 20 of narcolepsy. The term, "treatment" as used herein, refers not only to ameliorating symptoms associated with the disease, but also preventing or delaying the onset of the disease, and also lessening the severity or frequency of symptoms of the disease. In this embodiment, a nucleic acid of the invention (e.g., the HCRTR2 gene (SEQ ID NO:1)) can be used, either alone or in a pharmaceutical composition as described above. For example, the HCRTR2 gene, either by itself or included within a vector, 25 can be introduced into cells (either in vitro or in vivo) such that the cells produce native HCRTR2 receptor. If necessary, cells that have been transformed with the gene or can be introduced (or re-introduced) into an individual affected with the disease. Thus, cells which, in nature, lack native HCRTR2 expression and activity, or have mutant HCRTR2 expression and activity, can be engineered to express

-19-

HCRTR2 receptors (or, for example, an active fragment of the HCRTR2 receptor). In a preferred embodiment, nucleic acid comprising the HCRTR2 gene, can be introduced into an expression vector, such as a viral vector, and the vector can be introduced into appropriate cells which lack native HCRTR2 expression in an animal. In such methods, a cell population can be engineered to inducibly or constitutively express active HCRTR2 receptor. Other gene transfer systems, including viral and nonviral transfer systems, can be used. Alternatively, nonviral gene transfer methods, such as calcium phosphate coprecipitation, mechanical techniques (e.g., microinjection); membrane fusion-mediated transfer via liposomes; or direct DNA uptake, can also be used.

10

The nucleic acids and/or vectors are administered in a therapeutically effective amount (i.e., an amount that is sufficient to treat the disease, such as by ameliorating symptoms associated with the disease, preventing or delaying the onset of the disease, and/or also lessening the severity or frequency of symptoms of the disease). The amount which will be therapeutically effective in the treatment of a particular disorder or condition will depend on the nature of the disorder or condition, and can be determined by standard clinical techniques. In addition, in vitro or in vivo assays may optionally be employed to help identify optimal dosage ranges. The precise dose to be employed in the formulation will also depend on the route of administration, and the seriousness of the disease or disorder, and should be decided according to the judgment of a practitioner and each patient's circumstances. Effective doses may be extrapolated from dose-response curves derived from in vitro or animal model test systems.

The following Examples are offered for the purpose of illustrating the present invention and are not to be construed to limit the scope of this invention. The teachings of all references cited herein are hereby incorporated herein by reference.

-20-

EXAMPLES

EXAMPLE 1 Identification of the Human Narcolepsy Gene

A human BAC library (RPCI11 human male BAC library; see Osoegawa, K. et al., Genomics 52:1-8 (1998)) was used. Twenty primers, designed from the mRNA sequence of the HCRTR2 receptor, were employed to identify clones of interest. They are set forth in Table 1.

TABLE 1 Primers Used for Hybridization

	#	Name	Primer Sequence	SEQ ID NO:
	1	HCRTR2-1-F	TACTACTAGGCCACGCG	3
5	2	HCRTR2-1-R	ACACCAGGAGGAGAAAGCTAC	4
	3	HCRTR2-2-F	ATCGCCTGTAAAGACAGCAAAG	5
	4	HCRTR2-2-R	AAAGTTACTGAGCCAATGCCTC	6
	5	HCRTR2-3-F	GAGAGGAGCTTGCAGCATTG	7
	6	HCRTR2-3-R	AGGAATTCCTCGTCGTCATAGT	8
	7	HCRTR2-4-F	GAAGAACCACCACATGAGGAC	9
10	8	HCRTR2-4-R	ATCACTTTGCAAAGGGACTGTC	10
	9	HCRTR2-5-F	GTATGCAATCTGTCACCCTTTG	11
	10	HCRTR2-5-R	AATGCAGGAGACAATCCAGATG	12
	11	HCRTR2-6-F	CAGGCTTAGCCAATAAAACCAC	13
	12	HCRTR2-6-R	GATAAGCCAACACCATGAGACA	14
15	13	HCRTR2-7-F	ACAGATCCCTGGAACATCATCT	15
	14	HCRTR2-7-R	CTCGGATCTGCTTTATTTCAGC	16
	15	HCRTR2-8-F	CCAATTAGCATCCTCAATGTGC	17
	16	HCRTR2-8-R	GTGTGAAAAGGTAAACCAGGCA	18
	17	HCRTR2-9-F	CTCAGTGGAAAATTTCGAGAGG	19
20	18	HCRTR2-9-R	GTTGCTGATTTGAGTGGTCAAG	20 .
į	19	HCRTR2-10-F	CTTTCTGAGCAAGTTGTGCTCA	21
	20	HCRTR2-10-R	TACCAGTTTTGAAGTGGTCCTG	22

Initial Study with Large Membranes

Four out of 5 membranes having the whole BAC library, containing a total of approximately 160,000 BAC clones representing an approximately 10-fold coverage of the human genome, were used in hybridization studies with these primers.

Hybridization was performed with a pool of all 20 primers described in Table 1.

-22-

5' End Labeling for Big Membranes

Oligonucleotides were labeled at the 5' end before hybridization, using fresh (less than one month old) $[\gamma^{32}P]ATP$ (6000 Ci/mmole; 10 μ Ci/ μ I). The following protocol is adjusted for 4 membranes in 2 bottles, containing 2 membranes/30 ml of rapid hyb. Each. Briefly, a labeling mixture was made of DNA (8 pmol/µl) (10.0 µl of the primer pool), 10X buffer (12.0 μ l), T4 PNK (10 u/μ l) (6.0 μ l), [γ^{32} P]ATP (30.0 μ l, or 600 μ Ci), and water (62.0 μ l) for a final volume of 120 μ l. 20 μ l of labeling mixture was used per 10 ml rapid hybridization reaction. Incubation of the labeling mixture was for 2 hours at 37°C, followed by transfer to ice, spinning down, and mixing with the rapid hybridization solution. The membranes were prehybridized at 42°C before the labeling mix was added. Sixty µl of the labeling mix was added to each of 2 big bottles containing 2 membranes and 30 ml of rapid hybridization solution.

Hybridization and Washing

15

The membranes were hybridized at 42°C overnight. After overnight, membranes were washed with 6x SSC, 0.1% SDS at room temperature; washed with 6x SSC, 0.1% SDS at 55°C in a shaking waterbath, repeated until the radioactivity of membranes was lower than 6k using 1x sensitivity; and washed with 6x SSC to remove the SDS. The washed membranes were put in a cassette for overnight 20 exposure at -80°C with a MR single emulsion film. Positive clones were identified and gridded on small membranes.

Study of Positive Clones with Small Membranes

After growing the positively-identified clones on several small membranes (to get several copies of membranes containing the same clones), and washing the membranes, hybridization was performed using pairs of primers, instead of a total 25 pool of primers as before. The total number of hybridizations was ten, using different primers against identical copies of membranes containing all positive clones from the first hybridization. The primer pairs are set forth in Table 2; primer numbers indicate the primers shown in Table 1.

TABLE 2 Primer Pairs Used for Hybridization

	Reaction number	Primers Used
	1	1 and 2
	2	3 and 4
5	3	5 and 6
	4	7 and 8
	5	9 and 10
	6	11 and 12
	7	13 and 14
10	8	15 and 16
	9	17 and 18
	10	18 and 19

5' End Labeling for Small Membranes

Oligonucleotides were labeled at the 5' end before hybridization, using fresh

[γ³²P]ATP (5000 Ci/mmole; 10 μCi/μl). Briefly, a labeling mixture was made of DNA (8 pmol/μl) (1.5 μl), 10X buffer (2.0 μl), T4 PNK (10 u/μl) (1.0 μl),

[γ³²P]ATP (3.0 μl), and water (12.5 μl) for a final volume of 20 μl. Incubation of the labeling mixture was for 2.5 hours at 37°C, followed by transfer to ice, spinning down, and mixing with the rapid hybridization solution. Membranes were prewetted in 6X SSC, rolled in a pipette, and excess liquid drained prior to placing the membrane in the tube. Fifty ml Falcon (polypropylene) tubes were used as container for the hybridization. The membranes were prehybridized at 42°C before 20 μl of labeling mix was added to each tube.

Hybridization and Washing

The membranes were hybridized at 42°C overnight. After overnight, membranes were washed as described above. Four clones which were positive for primers designed using the 5' and 3' end of the mRNA were identified. Clone 403B19 was used to characterize the gene.

Sequencing of Narcolepsy Gene in Clone 403B19

Shotgun sequencing was used to obtain the gene sequence.

Preparation of DNA Samples

BAC DNA was isolated using the Plasmix kit from TALENT-VH Bio 5 Limited. Thirty μg of isolated DNA was fragmented by nebulization: a nebulizer (IPI Medical Products, Inc., no. 4207) was modified by removing the plastic cylinder drip ring, cutting off the outer rim of the cylinder, inverting it and placing it back into the nebulizer; the large hole in the top cover (where the mouth piece was attached) was sealed with a plastic stopper; the small hole was connected to a 1/4 inch length of Tycon tubing (connected to a compressed air source). A DNA sample was prepared containing 30 μg DNA, 10 X TM buffer (200 μl), sterile glycerol (1 ml), and sterile dd water (q.s.) for a total volume of 2 ml. The DNA sample was nebulized in an ice-water bath for 2 minutes and 40 seconds (pressure bar reading 0.5). The sample was then briefly centrifuged at 2500 rpm to collect the DNA; the entire unit was placed in the rotor bucked of a table top centrifuge (Beckman GPR tabletop centrifuge) fitted with pieces of Styrofoam to cushion the nebulizer. The sample was then distributed into four 1.5 ml microcentrifuge tubes and ethanol precipitated. The Dried DNA pellet was resuspended in 35 μ l of 1X TM buffer prior to proceeding with fragment end-repair. 20

Fragment End Repair, Size Selection and Phosphorylation

The DNA was resuspended in 27 μl of 1X TM buffer. The following materials were added: 10 X kinase buffer (5 μl), 10 mM rATP (5 μl), 0.25 mM dNTPs (7 μl), T4 polynucleotide kinase (1 μl (3 U/μl)), Klenow DNA polymerase (2 μl (5 U/μl)), T4 DNA polymerase (1 μl (3 U/μl)), for a total volume of 48 μl. The mixture was incubated at 37°C for 30 minutes, and then 5 μl of agarose gel loading dye was added. The mixture was then applied to separate wells of a 1% low melting temperature agarose gel and electrophoresed for 30-60 minutes at 100-120 mA. The DNA was then eluted from each sample lane, extracted from the agarose

using Ultrafree-DA columns (Millipore) and then cleaned with Microcon-100 columns (Amicon), precipitated in ethanol, and resuspended in 10 μ l of 10:0.1 TE buffer.

Ligation

EcoRV-linearized, CIAP-dephosphorylated Bluescript vector was used as a cloning vector. The following reagents were combined in a microcentrifuge tube, and incubated overnight at 4°C: DNA fragments (100-1000 ng), cloning vector (2 μl (10 ng/μl)), 10X ligation buffer (1 μl), T4 DNA ligase (NEB 202L) (1 μl (400 U/μl)), sterile dd water (q.s.), for a total of 10 μl.

10 Transformation of Ligated Products

The ligation products were diluted 1:5 with dd water and used to transform electrocompetent TOP 10F cells (Invitrogen) using GenePulser II (Biorad; voltage, 2.5 W, resistance 100 ohm). Transformants were plated on LB plates with 50 µl of 4% X-GAL and 50 µl of 4% IPTG, and ampicillin. Transformants were grown overnight at 37°C, white colonies were picked, grown in a culture of 3 ml LB liquid media plus 200 µg/µl ampicillin for 16-20 hours with shaking. DNA was isolated from the liquid cultures using Autogen 740 Automatic Plasmid Isolation System.

Cycle Sequencing of Isolated Plasmid DNA

Isolated plasmids were then sequenced using the M13 primers: M13-forward (SEQ ID NO:23) TGTAAAACGACGGCCAG; and M13-reverse (SEQ ID NO:24) CAGGAAACAGCTATGAC. For the sequencing reaction, 2.5 µl plasmid template was mixed with 4 µl Big Dye Ready reaction mix (ABI), 1 µl of 8 pM M13 primer, and 2.5 µl dd water. For cycle sequencing, 25 cycles of 96°C for 10 seconds, 50°C for 5 seconds, and 60 °C for 4 minutes were performed, followed by holding at 4°C.

The cycle sequencing reaction products were cleaned by spinning through Sephadex G-50 columns. The eluted cycle sequencing products were then dissolved in 3 µl formamide/dye and 1.5 µl of sample was loaded on ABI 377 automated sequencers. The data was analyzed using Phred and Phrap (Ewing, B. et al., Genome Res. 8:175-

10

20

185 (1998); Ewing, B. and Green, P., Genome Res. 8:186-194 (1998)), and viewed in Consed viewer (Gordon, D. et al., Genome Res. 8(3):195-202 (1998)).

Analysis of Gene Structure

The hcrtr-2 gene maps to chromosome 6p11-q11. A total of 168,575 base pairs of contiguous sequence was generated for 403B19 which contained all of the hcrtr-2 gene. Comparison of the cDNA sequence of hcrtr-2 (Accession number GI:6006037) and the genomic sequences generated allowed deduction of the intron/exon organization of the gene. The gene contains 7 exons which cover 108,439 bp. The first 10 Gs in the mRNA sequence for hcrtr-2 were not found in the genomic sequence. It is likely that these Gs were an artifact.

The splice junctions of the *hcrtr-2* gene are set forth in Table 3, and the intron sizes are set forth in Table 4. Exon sequences are represented in uppercase and introns in lowercase. All splice sites conform to the consensus GT-AG rule. SEQ ID NOs are given in the column immediately following each site.

15 Table 3 Splice Junctions of hcrtr-2

	Splice Donor Site	SEQ ID	Splice Acceptor Site	SEQ ID
Hcrtr-2 exon1-2	TCCTGGgtgagt	25	aattagTTTGTG	26
Hcrtr-2 exon2-3	CTACAGgtaatt	27	ctctagACCGTG	28
Hcrtr-2 exon3-4	GGGTGgtaagt	29	tcctagGTGAAA	30
Hcrtr-2 exon4-5	CGACAGgtatat	31	tttcagATCCCT	32
Hcrtr-2 exon5-6	AAAGAGgtaaaa	33	ctgcagAGTATT	34
Hcrtr-2 exon6-7	TCAGTGgtgagt	35	tgccagGAAAAT	36

5

Table 4 Intron Sizes of hcrtr-2

Intron	Nucleotides	
Intron 1	73,848	
Intron 2	6,322	
Intron 3	8,327	
Intron 4	13,618	
Intron 5	2,730	
Intron 6	1,779	

The exons do not clearly respect the domain structure of this seven

membrane domain G protein linked receptor. Five of the transmembrane regions are
by themselves within one exon, two of the transmebrane segments are broken up by
introns, and two transmembrane segments fall within the same exon. A survey done
one year ago on mammalian G-protein coupled receptors (GPCRs) sequences in
GenBank revealed that over 90% of GPCRs genes were intronless in their open

reading frame (ORF) (Gentles, A.J. and Karlin, S., Trends Genet. 15:47-49 (1999)).
Comparison of the intron/exon boundaries of hcrtr-2 and the genes coding for their
most related GPCRs based on sequence similarity showed that the location of the
intron/exons boundaries with respect to the transmembrane domains is only partially
conserved among the receptors (Sakurai, T. et al., Cell 92:573-585 (1998)).

20 Computer analysis of sequence data

Analysis of the genomic sequence of hcrtr-2 using the program RepeatMasker (http://ftp.genome.washington.edu/cgi-bin/RepeatMasker) showed that the sequence containing the hcrtr-2 genomic sequence is 38.27% repeat sequences and the GC content is 35.3%.

The sequences of the genes were analyzed using the program GeneMiner (Óskarsson and Pálsson, unpublished), which combines the results of 5 exon prediction programs; FGENE (Solovyev, V. and Salamov, A., *Ismb* 5:294-302 (1997)), Genscan (Burge, C. and Karlin, S., *J. Mol. Biol.* 268:78-94 (1997)),

30

HMMgene (Krogh, A., Ismb 5:179-186 (1997)), MZEF (Zhang, M.Q., Proc. Natl. Acad. Sci. USA 94:565-8 (1997)) and Xpound (Thomas, A. and Skolnick, M.H., IMA J. Math Appl. Med. Biol. 11:149-160 (1994)). For hcrtr-2, 3 out of 5 programs predicted the 3' end of exon 1, only one program predicted the 7th exon and for the internal exons, there were at least two programs that predicted each of them exactly or in part.

The promoter sequences of the genes have not yet been characterized. The Promoter Prediction by Neural Network

(http://www.fruitfly.org/seq_tools/promoter.html) predicted promoters that are at least

10 140 bp upstream of the 5'UTR of hcrtr-2, indicating that either a part of the 5'UTR is missing in the published mRNA sequence or the real promoters are not detected by the program.

Analysis of Population for Polymorphisms

in nucleic acid samples from 47 patients and 75 control individuals. The patient population consisted of patients of Icelandic and US origin. The control population consisted of Icelandic controls, CEPH (Centre d'Etude du Polymorphisme Humain) individuals from Utah and France, and US samples of various ethnic origins. The African-American/Caucasian ratios were similar between patients and controls. All narcoleptic subjects complained of excessive daytime sleepiness (EDS). Approximately 66% of the patients had cataplexy, 24% did not and 10% did not have attainable records of cataplexy status. Narcoleptic subjects without cataplexy had Multiple Sleep Latency Tests showing mean sleep latencies of less than 10 minutes and REM sleep in at least 2 naps. Subjects did not take any drugs affecting sleep for at least 10 days before their sleep studies.

To analyze the nucleic acids, DNA from patient and control blood samples were isolated by the method of Kunkel (Kunkel, L.M. et al., Proc. Natl. Acad. Sci. USA 74:1245-9 (1977)). Briefly, white blood cells were lysed in a sucrose lysis buffer, and proteinase K treated; the DNA was then extracted using phenol-chloroform/isoamylalcohol and then ethanol precipitated. Patient samples that were

-29-

received in the form of nuclei pelleted through sucrose buffer were resusupended in lysis buffer (100 mM NaCl2; 10 mM TrisHCl, pH 8; 25 mM EDTA pH 8; 0.5% sodium dodecyl sulfate; 0.1 mg/ml proteinase K) at 55°C for 4-6 hours followed by classical phenol-chloroform extraction and ethanol precipitation (Sambrook, J. et al., Molecular Cloning, A Laboratory Manual (1989)). Samples were incubated at 55°C after isolation for the inactivation of DNAse to prevent the degradation of DNA. Concentration of the isolated DNA was determined by spectrophotometric analysis at 260 nm (Sambrook et al., using GeneQuant (PharmaciaBiotech), and samples diluted with sterile distilled water to a 20 ng/µl working solution.

Primers were designed from intronic sequences flanking the exons of the hypocretin receptor-2 (*hcrtr-2*), using either primer design programs available at primer3 at the Whitehead Institute (http://www-genome.wi.mit.edu/cgi-bin/primer/primer3.cgi) or primers for the worldwide web (http://williamstone.com/primers/javascript/). The primers are shown in Table 5.

10

Table 5 Primers Used to Amplify Nucleic Acid Fragments for Analysis of hcrtr-2 Gene

	EX-	#	Primer Sequence	Sense/	External/	SEQ
	ON			Antisense	Nested	ID.
5	1	1	TTTCTTCAGCTTCAGCTCTCCCTCAGC	s	E	37
	1	2	TTCAGCTCCGAAGCAGATGACCAGTTG	A	E	38
	1	3	TTCAGCTTCAGCTCTCCCTCAGCGAGG	s	N	39
	1	4	CGAAGCAGATGACCAGTTGCGACAAGG	A	N	40
	1	5	CTTTCCCACCGCAAATCACCAGTGCTC	S	Е	41
10	1	6	ATTTTATTAGAAAACCCCATCCGAGAG	A	Е	42
	1	7	TTCCCACCGCAAATCACCAGTGCTC	s	N	43
	1	8	TATTAGAAAACCCCATCCGAGAGCAG	Α	N	44
	2	9	GCATGTACTTAGCATTCACACAGATTG	s	Е	45
	2	10	TCTAATGATGATTTGGCAGTTCATTGC	A	E	46
15	2	11	TAGCATTCACACAGATTGACAGATTCA	S	N	47
	2	12	CAGTITGTCAATGCCTTAGGCAAATAT	Α	N	48
	3	13	TTTGGCAGCTTTGAATTTGCTTATATG	S	E	49
	3	14	GCTCTTGCAAAACTGTATTCACAAATG	A	Е	50
	3	15	CAGCITTGAATTTGCTTATATGTTGTG	s	N	51
20	3	16	TTGCAAAACTGTATTCACAAATGTCAA	Α	N	52
	4	17	TCCCCTTTGCATACATAATATGACAATG	S	Е	53
	4	18	AAAAAGCACAGACAAAATATTTGGAAGG	A	E	54
	4	19	ATGCACTTTGAAGAAAAGCATTGACATG	S	N	55
	4	20	AAGCACAGACAAAATATTTGGAAGGAAT	Α	N	56
25	5	21	CTCAGGCGTCTGGAAGCCTTTCCTTAC	S	E	57
	5	22	TTAAAGGCTGTTCGCCTTACCTGCTGG	A	E	58
	5	23	GGCGTCTGGAAGCCTTTCCTTACTGTG	s	N	59
;	5	24	CTGAGTCATCTGGCCTGACAAGGTATC	Α	N	60
	6	25	GGGTCAGAAACCAATCTGTGGTCAATTC	S	E	61
30	6	26	AGTTGAAGAGTGTTCATTGATTCCTCATCC	A	E	62
	6	27	AGAAACCAATCTGTGGTCAATTCCTGCAAC	S	N	63

-31-

EX-	#	Primer Sequence	Sense/	External/	SEQ
ON			Antisense	Nested	ID.
6	28	TGAAGAGTGTTCATTGATTCCTCATCCTTG	A	N	64
7	29	GAGTCTACCAAGCTTCCAATAAACTCA	s	Е	65
7	30	GGATAGTTTTACTCAGGTATCCTTGTCA	A	Е	66
7	31	CAAATCAGCAACTTTGATAACATAT	s	N	67
7	32	GTATCCTTGTCATATGAATAAATATTCTAC	A	N	68
7	33	CACTCAAATCAGCAACTTTGATAAC	s	E	69
7	34	GTGAGAGATTAAAATAACAAGGGAT	A	Е	70
7	35	CAAATCAGCAACTTTGATAACATAT	s	N	71
7	36	TGTTTAAACATTTAATTGACACACA	A	N	72
7	37	TTCATATGACAAGGATACCTGAGTAAA	S	Е	73
7	38	GTGAAATAGCCTGAAATAAGCTCAA	A	Е	74

10

20

5

PCR reactions were done in 20 µl reactions using 40 ng genomic DNA, 0.2 mM solution of the four dNTPs, 0.35 µM of each primer (TAGCopenhagen), 2.5 mM MgCl2 (Perkin Elmer), 1x PCR Buffer (Perkin Elmer) and 0.5 U Amplitaq gold 15 (Perkin Elmer). The primers were used to amplify the fragments by PCR cycling at 95°C for 12 min and subsequently 30 cycles of 95°C for 30 sec, 55-62°C for 30 sec and 72°C for 1 min. The PCR products were prepared for cycle sequencing by incubation with Shrimp alkaline phosphatase (Amersham) and exonuclease I (Amersham) at 37°C for 15 min. After the inactivation of the enzymes the products were subject to cycle sequencing using BigDye Ready Reaction mix (Perkin Elmer) and subsequently run on ABI Prism 377 Automated DNA sequencers. The raw data were basecalled and sequences assembled using the Phred and Phrap software, respectively (Ewing, B. et al., Genome Res. 8:175-185 (1998); Ewing, B. and Green, P., Genome Res. 8:186-194 (1998)). The Consed viewer was used to analyze the sequences (Gordon, D. et al., Genome Res. 8(3):195-202 (1998)). Expansion of a T-25 stretch in the 3' untranslated region (UTR) of exon 7 of hcrtr-2 was investigated by amplifying a fragment containing the stretch with a fluorescently labelled primer

pair using the conditions described above. The PCR product was dissolved in formamide/dye solution and run on ABI Prism 377 Automated DNA sequencers as described above. Allele calling was done using TrueAllele and editing was done using DeCODE-GT (Palsson, B. et al., Genome Res. 9:1002-1012 (1999)).

A total of nine single nucleotide polymorphisms were identified, 7 in exons and 2 in an intronic sequence near an exon. The polymporphisms are shown in Table 6. The base number is according to the mRNA sequence (Accession number GI:6006037). For those polymorphisms marked with an asterisk (*), the polymorphism is located 5' of the corresponding exons; the numbers indicate the distance into the introns.

Table 6 Single Nucleotide Polymorphisms in hcrtr-2

5

10

	Location	cDNA base #	Nucleic Acid Change
	Exon 1	352	С-Т
	Exon 1	355	C-A
15	Intron1	-26*	C-A
	Exon 5	1,170	G-A
	Exon 5	1,177	C-A
	Exon 5	1,201	G-A
	Exon 5	1,246	G-A
20	Exon 5	1,266	G-A
	Intron 6	-87*	G-A

While this invention has been particularly shown and described with references to preferred embodiments thereof, it will be understood by those skilled in the art that various changes in form and details may be made therein without departing from the spirit and scope of the invention as defined by the appended claims.

CLAIMS

What is claimed is:

- Isolated nucleic acid molecule comprising the nucleic acid having SEQ ID
 NO:1.
 - A DNA construct comprising the isolated nucleic acid molecule of Claim 1
 operatively linked to a regulatory sequence.
 - A recombinant host cell comprising the isolated nucleic acid molecule of Claim 1 operatively linked to a regulatory sequence.
- 10 4. A pharmaceutical composition comprising a nucleic acid comprising the isolated nucleic acid molecule of Claim 1.
 - Isolated nucleic acid molecule comprising the nucleic acid having SEQ ID
 NO:1 with one or more of the nucleic acid changes shown in Table 6.
- 6. A method of diagnosing narcolepsy in an individual, comprising detecting a mutation in the gene encoding hypocretin (orexin) receptor 2, wherein the presence of the mutation in the gene is indicative of narcolepsy.
 - 7. A method of treating narcolepsy in an individual, comprising administering to the individual an isolated nucleic acid of Claim 1 in a therapeutically effective amount.

1/51

```
LOCUS
                                    168,575 bp DNA
                                                          PRI
                                                                 20-OCT-1999
DEFINITION
              Human hypocretin (orexin) receptor 2 (HCRTR2) gene, complete cds.
ACCESSION
NID
VERSION
KEYWORDS
SOURCE
              human.
 ORGANISM
              Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
              Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
              1 (bases 1-168,575)
AUTHORS
 TITLE
              Direct Submission
JOURNAL
                      Submitted (
                                         ) deCode Genetics, Inc., Lynghals 1,
                      IS-110 Reykjavik, Iceland.
FEATURES
                     Location/Qualifiers
                      1.. 168,575
       source
                     /organism="Homo sapiens"
                     /db_xref="taxon: 9606"
                     /chromosome="6"
                     /map="6p11-q11"
                     /clone="BAC 403B19"
                      1..129,305
       gene
                     /partial
                     /gene="HCRTR2"
                     /note="OX2R"
                     /db_xref="LocusID:3062"
                     /db_xref="MIM:602393"
                     20,867..21,403
      exon
                     /gene="HCRTR2"
                     /number=2
      CDS
                     join(21,181..21,403, 95,252..95,430, 101,753..101,996, 110,324..110,439,
                     124,058..124,278, 127,009..127,130, 128,910..129,139)
                     /gene="HCRTR2"
                     /note="HCRTR2 exons defined by comparison to mRNA sequence (NM_001526)"
                            /product="HCRTR2/orexin 2 receptor"
                     /db xref="LocusID:3062"
                     /db xref="MIM:602393"
                     /protein_id="NP_001517.1"
                     /db_xref="PID:g4557639"
                     /db_xref="GI:4557639"
                     translation="MSGTKLEDSPPCRNWSSASELNETQEPFLNPTDYDDEEFLRYLW/
                     REYLHPKEYEWVLIAGYIIVFVVALIGNVLVCVAVWKNHHMRTVTNYFIVNLSLADVL
                     VTITCLPATLVVDITETWFFGQSLCKVIPYLQTVSVSVSVLTLSCIALDRWYAICHPL
                     MFKSTAKRARNSIVIIWIVSCIIMIPQAIVMECSTVFPGLANKTTLFTVCDERWGGEI
                     YPKMYHICFFLVTYMAPLCLMVLAYLQIFRKLWCRQIPGTSSVVQRKWKPLQPVSQPR
                     GPGQPTKSRMSAVAAEIKQIRARRKTARMLMVVLLVFAICYLPISILNVLKRVFGMFA
                    HTEDRETVYAWFTFSHWLVYANSAANPIIYNFLSGKFREEFKAAFSCCCLGVHHRQED
                    RLTRGRTSTESRKSLTTQISNFDNISKLSEQVVLTSISTLPAANGAGPLQNW"
      exon
                    95,252..95,430
                    /gene="HCRTR2"
                    /number=3
                    101,753..101,996
      exon
                    /gene="HCRTR2"
                    /number=4
```

FIG. 1A

2/51

exon 110,324..110,439
/gene="HCRTR2"
/number=5
exon 124,058..124,278
/gene="HCRTR2"
/number=6
exon 127,009..127,130
/gene="HCRTR2"
/number=7
exon 128,910..129,305
/gene="HCRTR2"

BASE COUNT 55,308 a 29,672 c 29,838 g 53,757 t

/number=8

CGACTTGATTTTATTTTTTGCATATGGATATCCAGTTTTCACAGCACTGCTTGTTACCCT CAGCAAAGAACAGTTGTCTGTAAATTCATGGGTTTATGTCTAGGCTCTCTGTTCTGTTCT ATTGGTCAACATATGGTCATATATCACTTAACTGCAGGGAAGGGATACATTCTGAGAAAT GCATTATTACATGATTTCATCATTGTGCAAACACTATAGAGTGTAGTTACAGAAACCTAG TATCTCTAGCTGTTCTTATGATTCAAATTTGCTTTGGTCATTTGAGATCCATACTGGT GGAGTCTAATTATTCAAAACTAGGGAAAACAGACAAACAGAAAAAACTAAGACCAAGTTA CCCTTACCTTGACTAAAAAAAGAGACTCAAATAAATAAAATTGGAAATGAGACAGGAGAC ATTACAATTGATGTTAACAAAAAGATCATAAGGTACTATTATGAACAACTATACACCAAT AAATTGGACAACCTAGAAAAAAAATGGATAAATTCCTAGAAATACACAGTCTATCAAACT GAAACAAGAAGAAATAGAAAGCCTGAACATACCAGTAACAACCAAGGAGACTGAGTAAAT AATCAAAAACCTCCCAAGAAGAAGAGTCTAGGACCAGAAGTCTTCACAAATGAATTCTAC CAAACATTTAAAGTATTAATGCCAATCATTCATTCTTATACTCTTCCAAAAAGAAGAAGGG GAATATTTTCAAACTCATTTTATGAGGCCAGCATTATTCTGATACCAAAACTACGCAAAA ATACTACAAGAACATAAAAACTACAAATGTGGGAATTATCATGTATACATATGCAAAAAT CCTCAGTAAAATCCTAGCAAACTAAATTCAACAGTACATTAAAAAGATCATATAGCATGA CCAGTGAAATTTCTCCTTAGGACGCAAGGATAAGTCAACATATAAAATTGAATGTGATAT ACCACTTTAACAAAATGAAGGATAAAAATCATATGATCATCTGAATAGATGCAGAAAAAG CATATAACAAACTTTGACGTTGTTGAGAAATTGAAAGCTTTTCTCTAAGATCAAGAACAA AGCAAGGATGCCCATTCTTGCTTCTATTCAGCATAGTGCTTGAAGTCCTAGTCTGGACAA ATAAAATCCACCAAATTGGAAAGGAAGAAGTGAAATTACCTCTGTTTGTAGATGAGCTGA ATAGCTAGAGCAAAGAATGAATTCAGTACAGTTGCAGAATGCAAAATCAGTATACAAAA AGTACTTGTAATTCTATATAATAGCAACAAACTATTTCATAAGGAAATTAAGGAAACAAT CCCCATTACAATAGCATCATAAATAATAAATCTTAAGAACAAATTTAACCAAGGAGGTGA AAGACTTGTGTACTGAAAACTATAAAATGCTGATAAAAAAATTAAAGAAGATACAATAAA TGGAAGATATTCCATGCCATGGTTTGGAAGAATTAATATTGCTAAATGTACATACTACCC AAATGAAAACACAATCTTAACACTATTTAAACCAATTAAACAAACCTATGATTTCAATTT GGTCAAATGTGTTAGAATGGATTTCCTTTTATTGTTTTTGAACTTGTCTCTCCAAATTTC AAAGCCTGGTTCCTAATTTTTACTTGAAATACCAAATAACAAACCCACTTAATGAGCTCT GAGCCAGTTTTAGTAGCCAAACTTGTATTTAAATAGTGTGTTACATATTTGCACAAAAAG CCAACGGAGTCTAAATCAACACTAATTCACATCATTACTAGCAATCTAAAACATCAGATG ATAATTTTGCTGTTGTCTTTCAGGCAAGATATTCAACCATTGGTATTAAATGTTTTATAT GAATGTGCGGTGTTTTATTTCAGAAACACTTCTCTGAATTCCCAAGGCCTAAGAGCTATT CATCATAGAGGTTTGTGGAGGCGGTAGTTAGACATTTTCTACATGCATAATGTTAATTCA TTCAAACATTATAGAAAAAAGTTTGTAAAGAAGTTAATTTTCAAGGTGACAAAAAAATC AGATTGAATCATGTTTATTTTATTTCAATTTAAACTCGTTGGCTATCTTAGGAAATTCAC ATTGTTTTTGAAGAATATATGAACAAAGTTTGATTCATCTTATCTATATAAGCATGAGAG

FIG.1B

3/51

AAATACCTAGATGAGAGTGAAAAATGACTAATTTTGTGACCATTGTTATATCATAGATTA ACTTGTTCTCTACTTCTAAGCTGTGTGATCTTGAAAAGTCATCTAAACTTCAGGTAC CATCCTCACTTGCAAAATGAGGGGAAAAACCCCAGCACCTTTAATATGGTGTTATGTGGA TGAAATAAGTTAATATATTAAGTGCTTAGGTTTCATGCACTTTCTATATAGTATTAAT **AATATTATTGTTACATTATTATAGTTACATTATTATTTTTTATTAATATTATTGGAACATG** AATGGAATTGTTGTGGCTCATTTTAAAGATGCTGCAATGGAGACCAAGAGAAATTAAGTA TAATAATCCTAGTTAAGGTACAGCCATTTCTAATTACATTTTCCAACTGCTGCTTTTACT AGAACACATAATAATCTTATTTAATTATCACAACAATTTCTGTGGAGTTACTATTAATCC AGAGATGAAGAATCTAAAACTCTAAATTATCAAGCAACTATTCCAGCTTTAAACAACAGT AAAACTGGAATTAAAACTAGAGTTTCTTTATGAGGCCAGTATTACTCTTATACTAAAGCA CAAGCACACACACACACCAGATCAATATAACTTATGGATGTTGATGCAAAAATTTT CAAAAATTAATAGCAAATCGAATCCAGCAGTATTTTAAAGGACTATACACCATGGACAA ATGGGGTTTATTCCTGGGATATAAAGTTGGCTAAACTTAATGAAAATCAACCAGTGCAAT AAATAATAGTAATTAAAAAAACATAATTATCTCACTAGATGTACAAAAAATGACAAGATC CAACATGTTTTCAATATAAAAGCATTCCACAGACTAGGAATAGAAGGGAACTTCCTCAAC TTTACAAAGAACATCTACAACGAAACCACAGCTAACATCATATTTAATGGTGGAAGACTG AAATCTCTGAATATTTTCCCCTAAGATCAGAAAAAGACAAAGAGGTCTACTAATTCTATT GGAAGTAAAATTACCTCTGGCAGATGACATAATCTTATACATGGAGAACTCCTAGAGATT ACATACACACTCACAACTACCAGAGTTAATAAATGGGTTCTACAAAGTTGCAGGATACAA TATCAATTCTCAAAAAACACTTGTATTTCTATACACTAGCAAATAACTCTGAAAATGAAA TTACCAAGAATGTACAAGTTTTATGTACTGAAAAATAAAAAATGTCATTAAAAATAGTTA AAGAGAATCTAAGCACATTGCAGTTTTCTGACTCCAGGCCCGGGCTCTTGGATGGCATCT CTGGATCCACTCAGGACCAGGGAGAACTTGTTGCCCTGAAGGGAAGGACACAAGTCTGAC TGGCTTTACCACCTGCTGATTGTAGAAGCCTAGGGCCTTCAGGGAACACAGGTGGTAGCC AGATAGCAGTTACCATGGGCATTAGGCATGACCCAGTGCTATGTTGGCTTCTAGTTTGAC CCAGCACAGCCCAAGGGTGGTAACCACATGGGTGCTTGTGTCACCCCTCCTTAAGTTCCA GGCAGCCAGCAAAGAGAGAGTGACTCTGTTTGGGAGAAAGTAAGGGAAGAAAAAAGT CTCTGTTGGTAATACAAGGAATTCTTCCAGATCTTATCCAAGACCTCTATGAATCTGCAA CAGCCAAAGCATTATTAGTTTTCAGGTTTCCCCAGTGCAGATATGACTGCAATGATCAAA GCACAAACAAGTTTGGACTGGGAGGACTACAATAAATACCTAACTTCTCAATGCCCAGAA AGGAACAGCTCTGGTCTGCAGCTCCCAGTGAGATCAGTGCAGAAGGTGGTGGTTTCTGCA CAGAGGGTGAGCCAAAGCTGGATGGGGTGTCACCTCACTGGGGAAGCACAAGGGGATGGG GAACTCCCTCCCCTAGCCAACGGAATCTGTGAGGGACTGCCATGAGGGATGGTGCATTCT GGTCCAGATACTATGCTTTTCCCATGTTCTTCACAACCCTCAGGCCAGGAGATTCCCTCG GGTGCCTACACCACCAGGGCCTTGGGTTTCAAGTACAAAACTGGGTGGATCTTTGGGCAG GCACCGAGCTAGCTGCAGGAGTTATTTTTCATACCCCAGTGGTGCCTGGAATGCCAGTGA GACAGAACCATTCACTCTCGGAAAGGGAGCTGAAGCCAGGGAACCCAGTGGTCTAGCT CGGTGGATCCCACTCCCATGGAGGCCAGTAAGCTAAGATCCACTGGCTTGAAATTCTCAC TGCCAGTGCAGCAGTCTGAAGTCAACCTGGGATGCTTGAGCTTGGTGGAGAGAGGGACGT CCACCATTACTGAGGTTTGAGTAAGCAGTTTTCCCCTCACAGTGTAAACAAAGCCACTGG GAAGTTAAAGTAGGTGGAGCCCACGACAGTTCGGCAAAGCCACTATAGCCAGAATGCCTC AGCTTATAGATCAAACTCCCATCTCCCTGGGGACAGGGCACCTGGGGAAAGGGGCCAGCTGT GGGTGCAGCTTCAGCAGACTTAAATATTGCCGCAAGCTGACTCTGAAGACAGCAGGGGAT CTCCCAGCACAGCGCTCGAGCTCTGCTAAGGGGCAGACTGCCTCCAAGTGGGTCTCTG GCAGGCAGCAATCTTTGCAGTACTGTAGCCTCTACTGGTGATACCCAGGCAAATAGGGTC TGACGTTGACCTCCAGCAAACTCCAGCAGACCTTCAGCAGACGGGCCTGAGTGTAAGAAG

FIG. 1C

ACAAAAACAAAAACAGCACATCCGCACAAAAACCCCCATCTGAAGGTCACCAACACCAAAT ACCAAAGGTAGATAAATCCACAAAGATGGGGAAAAACCAGCACAAAAAAGCTGAAAATTC CAAAAAACAGAATACCTCTTCTCCTCCAAAGGATCACAATTCCTCACCAGCAAGGGGACA AAACTGGACAGAGAATGAGTTTGATGAATTGACAGAAGTAGGCTTGAAAAGGTGGGTAAT AAACTCCTCTGAGCTAAAGGAGCATGTTCTAACCCAATGCAAGGAAGCTAAGAACCTTGA AAAATGGTTAGAGTAATTGCTAACTAGAATAACCAGTTTAGAGAAGAGCATAAATGACCT GATGGAGCTGAAAACTATAGCACAAGAACTTCGTGCAGCATACACAGGTATCAATATCCA AATCGATCAAGCAAAGAAAAGAATATCAGAGATTGAAGATCAACTTAATGAAATAAAGTG TGAAGACCAGATTAGAGAAAAAAGAATAAAAAGGAATGAACAAAGTCTCCAAGAAATATG GGAATATGTGAAAAGACTAAACCTACATTTGATTAGTGTACCTGAAAGTGACGGGGAGAA AGGAATCAAGTTGGAAAACATTCTTCAGGATATTATCCAGGAGAACATCCACAACCTAGC AAGACAGGCCAACATTTAAATTCAGGAAATACAGAGTACATCACAAAGATACTCCTCGAG AAAAACAACCCCAAGACACATAATTGTCAGATGCACCAAGGTTGAAATACAGGAAAAAAG TTAAGGGCAGCCAGAGAGAAAGGTCGGGTTACCCACAAAGGGAAGCCCATCAGACTAACA GTGGATCTCCCTGCAGAAACCCTACAAGCCAGAAGAGAGTGAAGGCCAATATTCAACATG CTTTAAGAAAAGAATTTCAACCCACAATTTCATATCCAGCCAAACTATGCTTCATAGTG AAGGAGAAATAAAATCCTTTACAGACAAGCAAATGCTGAGAAATTTTGTCACCACCAGGC CTGCCTTACAAGAGCTCCCGAAGGAAGCACTAAATATGAAAAGGAAAAACCAGTATCAGC CACTGCAAAAACATATGAAATTGTAAAGACCATCAACACTATGAAGAAACTGCATCAACT AATGGGCAAAATAACCAGCTAGCATTATAATGACAGGATCAAATTCACACATAACGATAT TAACCTTAAATGTAATAGGCTAACTGCCCCAATTAAGAGACACAGACTGGCAAATTGGAT AGAGAGTCAAGACCCAACAGTGTGCTGTATTCAGGAGTCCAATTCATGTGCAAAGATACA TATAGGCTCGAAATAAAGGGATGGAGGAATATTTACTAAGCAAATGGAAAGCAAAATAAA GCGGAGGTTGCAATCCTAGTCTCTGATAAAATAGACTTCAAAACCAACAAAGATCAAAAGA CTATAGAGACTTAGACTCCCACGTAATAATAGTGGGAGACTTTAACACCCCACTGTCAAT ATTAAACAGATCAATGAGACAGAAAATTAACAAGTACATTCAGGACTTGAACTCAGCTCT ATTATTCTCAGCACCACTTGCACTTATTCTAAAATTGACCACATCATTGGAAGTAAAAG AATCAAATAAGAGCTCTGGAATAAGAAACTCACTCAAAACCGCACAACTACATGGAAACT GTTACTTGAAACCAATGAGAACAAAGACACAACATACCAGAATCTCTGGGACACAGCTAA AGTAGTGTTTGGAGGGAAATTCATAGCACTAAATGCCCACACGAGAAAGTGGGAAAGATC ATAGATAGATCACTAGCCAGACTAATGAAGAAGAAAAGAGAAGAATTGTATAGACACA ATAAAAATGATAAAGGGGAGATCATCACTGATCCCACAGAAATACAAACTACCATCAGA GAATACTATAGACACCTCTATGCAAATAAACTAGAAAACCTAGAAGAAATGGATAAATTC CTGGACACATACACCTTCCCAAGACTAAACCAGGAAGAAGTCAAATCCCTGAACAGACCA ATAACAAGTCCTGAAATTGAGGCAGTAATTAATAGCGTTCCAATGAAAAAAAGCCCAGGA CCAGATGGATTCACAGCCAAATTCTACAAGAGGTACAAATCAGAGCTGGTACCATTCCTT CTGAAACTATTCCAAACAACAGAAAAGAAAGACTCCTCCCTAACTCATTTTATGAGGCT GGCATCATCCTGATACCAAAACCTGGCAGAGACATACACACAAAAAAAGAAAATTTCAGGC TAATATATCCCTGATTAACACCGACGCAAAAATCCTCAATAAAATACTGGCAAACCAAAT CCAGCAGCACATCAAAAAGCTTATCCACCACGATCAAGTTGGCTTCATACCTGGCATGCA AGGCTTGTTCAACATACGAAAATCAATAAATGTAATTCATCACAAAAACAGAACCAATGA CAAAAACCACATGATTATCTCAATAGATGCAGAAAAGGCCTTCAACAAAATTTAACAGCC CTTCATGCTAAAAACTCTCAATAAGCTAGGTATCGATGCAATGTATTTTAAAACAATAAG AGCTATTTATGACAAACCCATACCCAATATCATACTGAATGGGCAAAAGCTGGAAGCATT CCCTTTAAAAACTGGCACAAGACAAGGATGCCCTCTCTCACCACTCCTATTCAACATAGT AAGAGAAGTCAAATTGTCTCTGTTTGTGGATGACATCATTGTATATTTAGAAAACCC CATTGTCTCAGCCCAAAATCTCCTTAAGCTGATAAGCAACTTCAGCAAAGTCTCAGGATA CAAAATCAATGTGCAAAAATCACAAGCATTTCTATACACTAATAATAGACAAACAGAGAG

FIG. 1D

CCAAATCATGAGTGAACTCCCATTCAAAATACCTAGGAATACAACTTACAAGGGATGTGA ATGCAAAAACATTCCATCCTCATGGATAGGAAGAATCAATATCATGACAATGGCCATACT GCCCAAAATAATTTATAGACTCAATGCTATGTTCATCAAGCTACCACCGAATTTCTTCAC AGAATTAGTAAAAAACTGGCCAGGCTCAGTGGCTCACGCTTGTAATCCAAGCACTTTGGG AGGCCAAGGCAGGAGGATCAAGAGGTCAGGAGATTGAGACCATGGTGAAACCCCGTCTCT ACTAAAAATACAAAAAATTAGCCGGGCGTGGTGGCAGGCGCCTGTAGTCCCAGCTACTTG GAGAGGCTGAGGCAGGAGATGGCGTGAACCCAGGAGACGGAGCTTGCAATGAGCCAAGA ACAAACAACAAAAAAAAAAAACTACCTTAAATTTCTTATGGAACTAAAAAAGAGCCCAT ATAGCCAAAACAATCCTAAGCAAAAAGAACATAGCTGGAGGCATCATGCTACCTAACTTC AAATTATGCTACAAGGCTACAGTAACCAAAACAGCATGGTATTGGTATGAAAACAGATAT ATAGACCAATGGAACAGAACAGAGGCCTCAGAAATAACCCCAGACATCTACAACTCTCTG ATTTTTGACAAACCTGACAAAAACAAGCAATGGGGAAAGGATTTCCTATTTAATAAATGT TGTTGCGAAAACTGGCTAGCCATATGCAGAAAACTGAAACGGGACTCCTCCCTTACACCT TATACAAAATTAACTCAAGATGGATTAAAGACTTAAACGTAAGACCTAAAAACCATAAG AACCCTAGAAGAAAACCTAGGAAATACCATTCAGGCCATAGGCATGGGCAAACACTTCAT GTCTAAAACATCAAAAGCAATGGCAAGAAAATCCCAAATTGACAAATGGGATCTAATTAA ACTAAAGAGCTTCTGCACAGCAAAAGAAACTATCATCAGAGTGAACAGGCAACCTATAAA ATGGGAGAAAATTTTTGCAATCTGTCCATCTGATAAAGGGCTAATATCCAGAATCTACAA TGAACTCCAACAAATTTACAAGAAAAAAACAACCCCATCAAAAAGTGGGTGAAGGATGTG TCATCACTGGTCATTGGAGAAATGCAAATAAAAACCACAGTGAGATACCATCTCACTCCA GTTAGAATGGCGATCATTAAAAAGTCAGGAAACAACAGATGCTGGAGAGGATGTGGAGAA ATAGGAACGCTTTTACACTGTTGGTGGGAGTGTAAATTAGTTCAACCATTGTGGAAGACA GTGTGGTGATTCCTCAAGGATCTAGAACCAGAAATACCATTTGACCTAGCAATCCCATTA CTGGGCATATACCCAAAGGATTATAAATCATTCTATGATAAACACACATGCACATGTATG ATAGACTAGATTAAGAAAATGTGGCACATATACACCATGAAATACTATGCAGCCATAAAA AAGGATGAGTTCATGTCCTTTGCAGTGACATGAATGAAGCTGGAAACCATCATTCTCAGC ACAATGAGAACACATGGACACAGGGGGGGGGGAACATCACACACCAGGGCCTGTCAGGCAGT GGGGGCTAGGGGAGGGATAACATTAGGAGAAATACATAATGTAGGTGACAGGTTGATGG GTGCAGCAAACCACCGTGGCACATGTATACCTATGTAACAAACCTGCACGTTCTGCACAT GTATCCCAGAACTTAAAGTATTAAAAAAAAAAAAGACCATTTATGAAAACATGACCTTACCA AAGAACTATATAAGTCACTGGAGACCAATCCTGGAGTGACAGAAATATGTGACCTCTCAG ATGGAGAATTCAAAATAGCTGTTGTGAGGAAATTCAACAAAATTCAAGATGACATGGCAA AGGAATTCAGACTTCTATCAGATAAATTCAAAAAAGAAGATGAAATAATTTTTTTAAAAA TTCATGCAGAAATTTTGGAGCTGAAAAATTCAATTGATATACAAAAGAATGCATCTTACC AGCAGAATTGATCCTGCAGAAGAAAGAATTAGTAAATTTGAAAACACTCTATTTGAAAAT ATACAGTCAGAGGAGACAAAAGAAGAAAATTAAAAACAATGAAGCATACCTACAGGATCT AGAGAGAGAGTGGGATAGGGGTAGAAAGTTTATTCAAAGGGATAACAATAGAGTATCAGT ATTCAAATACAAGGTTATGGAACACCATTCAGATTTAACCCAAAGAAGACTACCTCAAGA CATTTAATAACTGAACTCTCATTCAATGGGAAAAGTAAAGTCCTTTCAATAAAGGTGTTG GGATAATTGGGTATGCAAAAAATGAATTTGGATACCTTTCTTGTGTCATATACATAAAAC CACAGTAAATTTTTGTGACCTTTGATTAGGCAATGATTTCTTAAATATGATAAAATATGG TAAAAGCAACAAAAGAAAACATGAATAAATTGGATCTTATCAAAATTTAAAACTTTTTTG CATCGTAGAATACTATCAAGAGTATGAAAAGAAAACCTACAAAATAGGAGAACATGTTTG TATATATATATATATATATACTCTTACACCTCAACTATAAAGAGACGAATAACCCAAT AAGTTCATCAAAAGATGCTCATCATCTTTACTCAGGAGGCAAATACAGATTAATATTACA ATGATATTAGACATGGATTTGTCATATACAGACTTTATTAAGTTAGATTCCCTCTATGCC TAATTTGTTGAGAGTTTTTATCATGAAGAGATGTTGCATTTTGTCAAATGCCTTTTCTGT GTCTTTTGAGATGATCATATGGTTTTCGTCCTTTATTTTGCTGATATGATGTACCACATT

FIG. 1E

TATTGATTTGCATTTATTGAATCATCCTTCCACCCCTGGGATAAATCCCACTTGATCATG GTGTATTATCTTTTTGATGTTTTTTGGATTCACTTTGCTGATATTTTGTTGAGGATTTCT GCATCTATAATCATTAAGGATATTGGCCTGTAGTTTTCTGTTTTTTATGTTGTATTCTAGT CTGATTTTGGTATCAGGGTAATGCTGTTCTTGTTGAGCGTGTCAGGAAGTCCAAAAGACT TAGAATTCAGCAGTAAAGCCATCCAGTTCTGGGCTTTTCTTTGTTAAGAGACTTAAAACA CACACAACGCACACAAAATGAAATATCACTTTCCACCCATTATAATTTACAAAGTGGA AAATAACTCGTGTTGATAAGAATGTGGAAACCTTGAAACCTTCATGCATTGCCAGTGGTA ATGTGAAAGAATCTTGCCATTGTGGAAAACAATTTGTCAGTTCCTCAAACAGTTCAACAT AGAGTTACTGTATGAAATAATTCAACTCCCAGGCATGCACCCAAGAGCATTGAAAACATA ATGGAAACAATCCAAATATTCATCAACTGCTGAATAGATAAAATGTGGCATATCCATATA ATTAAATACTATTCAGCCACAAAAATAATAAAGTACGGATAGACACTAAAACATGGAAGA ACCTTGAAAATATTAAGCTAAGTGAAAGACATAAGACACAAAACCCAACATTTAAAGGAA ATTTCCAGAATTGTCAGATCCACTGAAGAAGAAACTTGAGTGTTTGCCAGCATGTGGGAG GAGAGGAAAATCAGTAGTTATGAGGTTTCTGGAATTAGTAGTGCTGATGGTGACACAACA TTGTGAATATACTATAAACCACTAAATGATACCTCTCAAAATGGTTAAAACATTACTGTT GTGTTATGTGAATTTACCTCAATTAGAAAAGAAAAAATCTTATCAATAACAAAGAGAA ATTTCCACACAAGGTGGGATCGCTTCCACAGTGCTACTCAATGCAGTTTAGCGATTGCAT TTGTATTGGAGTAAAAGCATGTCACATTGCTTTTAACATTGGAGTCCAATACATAAACCT AATTTCACAACACAATGGATCATTTTTTTTTTTCATGTGGAAAATCAGAACACATGCCTTA ATGGTTACATGCCCCACCTGCTGCTCACCTAAAAGTAAATTTCCTCTAACTCAGACAAAT ATGTTATTTTCAAGGAAAAGAAGCCCAGAGAACTGAGATCCAGAAGAAATAACATGTATT GAAAGCACACAGAAGTATTTCAATGAACTCAAACCCAAGATTGTAGAAAACTCTCATGTG CCCACCCTTCAGAGCACCCGACGATAATGGATAGTTTCTAGCAGGGTGTCTGGAATGGGC AAGTACCCCCAAAGTTATAGTTTGTACTGCAAGACTTGAACCCACTCTTTTTCTGCCCTC TATTATTATTTTGCATTTTAACCATTTATTATTTTGAAAAGAAAAGAGAATTTTTAGAA TATGGAAAGAGGAAGTGAATTAATAAAATAGCACACCCTACATAGAGACTGCTAATCCAT CTCCAGTCTAAAGATTTAGTAATAGGCAAGAATATACATATCCAGGAATTTCCTTGGTGT TACATAAACAAAGGCGGCACATATGTATATTTTTCACAAAATATTCACTGTTTGAAGAAG GAATTACTCCCTTCAATTGAGTTCAGGCCTGATCAACAAGTAGTGATTGGCCAACAGCTA GGAGATGTCATCCTGAAGAGTATAACAAGTTCCCCTATAATTCTACTTTTCAGTACTGTT TAAAATACAACTGGATTTTTTTAAATATGTAAAATTTATATAATTTTACAAATGTCTTTG TTAAGAATTAAAACTATCATTAGTAAAGGACACAGCTGGAAAATTGAAAACATTTTGGTT CTCTACTGTGGAAACAGAATAGAGTAACAGCAAAAAGCGTATTTCTGGAATTGGACCCTG ACAACTCTGCTTAAACACTCCACCACTTTCTAGCTATATGACCTTGGGTAAGTTACTTAA CTTCTTTGTGTGTCAGTTTCTTCATTTGTAAAATTGGAATAATAGATGCTTTTTTTGAGA CAGTGTCTCATTCTGTTGCCCAGGCTGGAGTGCAGTGGCGTGACCACAGCTCACTGCAGC CTCAACCTCCTGAGTTCAAGTGATTCTCCAACTTGAGCCTCCCAGATAGCTAGGACCACA GACACATGCCACCATGCCTGGGTAATTTTTTTTTTAAGTTTTTCATAGAAATAGTGTCTC ACTAAGTTGCCCAACCTGGAAAATTGGAATAATAATTCATAAAATCTTCCTCCTAGATTT GTGAAGATCAATTGAGTTAATGTATGTAACGTACTTGGCACAGAGCTTGGCCCATGTAAT CAATGTCTTTTCCATATGGTTTCATTGACGCCACTTTGGGAAAATAGATGTCTCTTCTGC TTGCATTTTCAGACCTTTTTAGGTGTATACCTTAGGGCATTTGCTTTACTGACCAAAATT ATTTGCCGGCTACTCTGTGCTTTTCATGACACACTGAATAAGACAGGAAGAGTGTTTATC TATGCTCAACATAAGATAGGCATATAATGGAAGCTTCGTATATATTTGTTGAATAAAAAA CATAAGGGGAAAATATCAGATCTAATAATGCAGGACAGGAGGCAAGATGGAACGGAGAGA ACCTTGTCTGAGAAGAGACATAATTAAAACAGGGCATGGGAGGTAATAGAAAGATTGGAG GAAAAAGAGACAGAGAGACAGAAATGTTTGTGGTAATTTGTGACAAGTAGCTTTGATTGT TCATGGCCTAATCTTTTAGGGCATGAGGTTATTTCATTCTCTGTAGCCCACCGAGAGTGC GTACAGTGACACATGTTATGTAAGTCCCCTTTTTCCCTTTTTATAAATGTCTAGACCCCCT

FIG. 1F

7/51

GTGATTTGAGACTTTTCTAGAAGAATTTAGCTGAAGACCATATTGTTTTTTAAATGTAGT ATATAAGCTCAGTATCATTACCAACAGTGCTCAGACTTGATTTTATTTTCATTCCAA CAGCAAAGGAAAGAAAGCAACTTCTTTCATGCTTCCATGCCACCTCTGCATCTCTACCT TCACAGAGTTTCTCAATAATGGCAACATTTCCAGTTCACCAATGGACTGAGAGATCATTG **AGGCTAGACTAGTCTTATTAATCCTTATACCCCAGCTCCTAGCCGAACTCCTGGACACAC AATAGATACTCAGATACATTTACTGAAATGCATATAGAAAGTTACACCTGCAAAAAAGAT** GATCTCTCACCAGGAATAAGAAAATATAATCTGGGACAGCCCATATATGAGATCTCTAAA CTAGGTATTTAAACAGAATTATTCTGAATGTTGTGAGCTACATTTCTTTTTTACCTTTTA CATAGTATTTGTATATTTTATAGGGTACATGTAATATTTTGTTACACGCATAGAATGTG GCTAGGAGCATGTTAAGTCCTCTTTTTAGCTATTTTGAAATGTACATTGATGTTAACTA TCATTAACACAGAGTAATTGATATGTATAGCAAATAATATTTGCAGTAGGATATCACATG TTTACTTATTTATTTATTTATTTATTTATTATTATACTTTAAGTTCTAGGGTACATGTGCA CAACGTGCAGGTTTGTTACATATGTATGCATGCGCCATGTTGGTGTGCTGCACCCATTAA ACAGGTTCCAGTGTGTGATGTTCCCCTTCCTGTGTCCAGGTGTTCTCATTGTTTAATTCC CACCTATGAGTGAGAACATACGGTGTTTGGTTTTTTTGTCCTTGCGATAGTTTGCTGAGAA TCATGGTTTCCAGCTTCATCCATGTCTCTGCAAAGGACATGAACTCATCCTTTTTTTGGC TGCATAGTATTCCATGGTGTATATGTGCCATATTTTCTTAATCCAGTCTATCATTGTTGG ACATTTGGGTTGGTTCCAAGTCTTTGCTATTGTGAATAGTGCCGCAATAAACATATGTGT GCATGTGTCTTTATAGCAGCATGATTTATAATCCTTTGTGTATATACCCAGTAATGGGAT GGCTGGGTCAAATGGTATTTCTAGTTCTAGATCCTTGAGGAATTGCCACACTCTCTTCCA CAATGATTGAACTAGTTTACACTCCCACCAACAGTGCAAAAGTGTTCCTATTTCTCCACA TCCTCTCCAGCACCTGTTGTTTCCTGACTTTTTAATGATCGCCATTCTAACTGGAGTGAG GCACTGGTCTGAAAATATCAATTCATTTAATTCTTTTAACAACCTTAAGGGGATATCATG CAGCAGAAATATTTGAATTGAAGAGAAGAGTAATACCTAAGAACTAGAAATTCCTTTCTT ATGTTTCAAAAGATATCAAAAGATCTAAGGAAGATATTCACATCAAAAATGAGTATTATA ATATTATTATCTATGGTGCACTTGCAAAAAAGAAAACAAGTAATAATCTGAAGATTTAA GTGAATATTTTATGACATTGGAGTACCACATATTTAGAAGAAAGCACCAGAGAAATCATA GATAGAAGGAAATGGAATATTTGTAGGATCAAGATAAATACAGCTTGTCATAAAATAAAG ACTGTGATGATTAATTGTAGGTGGAAGATTTACGAAGAGAAGACTGAAGTATAGACAAGT TGAAGTGCCACAAAATGAAAGCTAATGACACTGACTACTTAGGAAATAGCAGACTGGGTC CATATTTATAGATTGTCAATGACAAGGAATTTGCAGATGTTAATGAATATAGATCCGAAC TTAAGTTGCAACAACCTTTCCCACTTTGAGATGAATAGTGCATGGAAGAGTAAAATGCAG ATGTTAATAAATCAGAGGAAGACATCGTGCCAGAGTATAAAGTTGACAGATTTATGCCGA ${\tt TGAACTTGAACAAAGCCACAGAAGGCCTACTTGTCAAATTTACTGGTGACAACAGGTCTG}$ GAGAAATGGCTAATGTTTTGGATAATAGCATTAGAATTTAAGGTCTGTTTAAACTTCAAA TTAACAGAATGAAATTAATATATGCACATATCAATTGGGTCTTTTGCTTATATATCATCT CTTAATAGAGCCTTTTTGAACAATCATTTCTAATGTGACCTTTGGGATTTTCTACTCATC ATCACCTCATCCTGTTTGGTTTGCATTATAGCATCTATCCCTTCCTAACGTTTTCCCTAT GTATTTGTTAGTTTTTTTTTTAATCTAACTTTACTAGAAAGTAAAATGCATGGAAAC AGCAACCTGTTTAACTTTGTATCACTAAGAGTGGAAAAATAACCCTCAGGAAATATTTGG TAAAATAATAAAATGCCCATTGATGCCCTTCTCTTAAAAAGAAATTTAATTAGTGCAGAT TGGGGAAATACAACAATATTTCTCATAAAATGTGATATCTATACAATAACAGAAGTACTA TGTCCCAAAAGTATTCTATAAATAGAAGAAGAACAGATGGTTTTGCTGCTGATTAATC CATTTATCTTTCGTAAATCATCTAATTTCCCCAGGAACAGCTTCCTCATCTATTAAAGGG GGTTAGTAATAGCTAAGCCCTCAGGGGTTTAAAAATGCATATGAAATAATTTTATAAACC ATAAAGCACAAAACAAATATGAAAAATTATGATTGGAGGAGGGGGTGGGGTAGTTAACTA AATCTCAGTGTAAACCACCAATGTCTTGTGTGTGTTGAAAAAATAATTACATATAAAAAC TGGTTGCATCCAAAGAATAATGTACTTTTTGCACTGGCAAGACTCAAACCATATTATTGT

FIG. 1G

TACTTCCTCCCAGTTACATATTTTGCAAGATATTGACAATTGTCTAAAGGAAGACCAAAC AACATGAGGGCATATAAAGGAGCGCTGGGGCTGTGATGTTTATTTTGAATCTGTGAAGCA TTGTCATGTGGAAGATTTATTCTGTGTAGCACCAAGATGCAAACTAGGAATTAGAGGTAA AAGTCTCAAAAAGACAAATCGTGGCTTGAGACCTTGGTTTAATGTAAGAAACAGTTTTCT CACCCTTAGAGCACTCCCATAAGGATGGAAGTAGTGAATTGTGGTGGTCACATTCAAGCT AGATGGGGACATGTCAGCAATGTTATCAGGAGGCTTCTACTCTGAAGCTGAAGTTCAGAC AAGATTTCCAGGCTCTTCCCAAGTGCAAGATTGTAATTACTTAAATGCAATATTTTTACC ATGTTTATTAAGAATAAAAGGATCATGAATTCACATTCTGACAAATGCTAGAATACTTAT TATTAGAGACAAAACCAGTGCATGAGAGAATGGCAGGTGACATCAGCCCTGAATCAATGG CATCCTACAGTGAATGTTTAATATCATTGAGTATATTGGTGGTCTGTCATGCTTGACAAC ATTAACTATGATCATATTTATGACACTTGGCGTCCTTCAAGAATTTGTAGCTCTATTTCA CATGACACTTAACTATCGCAAATACAAATTCCAGCTAAATAGACCCTTCAGTTTAAAAAC AGTCTCATTCTCAAATTTTAAGGAGAAAGTGAAGACGGAGATGTCTTAAAGACTCGGCAA GTACTAAGTTGGCAAATGTCAAATGTTAAAATAAGTTTATATTAAATGTTAAAGTGTTTG CCTGGAATGACTTTTCCATTGTCCTGCTTGAGAAACACAGAGGCACCTCCTTATTGCTTT TATATTTGCTTTACAAAGACAAATGTATCAACATGCTCTGTATTAATTGTATGTTGACAT TTTTGTCATATCCACAGACTGATGCATGTCTGTGCATGGTTTATAATAAGTGCACGTAAA AATAGAGAAAATAAGTAGAAAAAGAGAGAGATTTAACTCTCACCCCCCACCCCCAAAAA CGAGGGAGGAGGCTGTGGGCTGCGGACTGAGTGCTGGAATGAGGAGTAATTGAGCTTCAG CTGAGCCGGACGTAGCTTTCTCCTCCTGGTGTCATTGCTGCAGCCTCCAGTGCCGGGTCC CTAGTTCCTCAGCTGCCTATCTTCCCGGTGCAACATCGCCTGTAAAGACAGCCAACACCCAC $\tt CGCAGAAGTTGCCCGGCAGAAGACTCCGGAGGCATTGGCTCAGTAACTTTTCACGTCATT$ TTCTGCTCGGGAGCCCTTCTAGCCTCTCCGCGCAGCCTTTCCCACCGCAAATCACCAGT ATGTCCGGCACCAAATTGGAGGACTCCCCCCTTGTCGCAACTGGTCATCTGCTTCGGAG CTGAATGAAACTCAAGAGCCCTTTTTAAACCCCACCGACTATGACGACGAGGAATTCCTG CGGTACCTGTGGAGGGAATACCTGCACCCGAAAGAATATGAGTGGGTCCTGATCGCCGGG TACATCATCGTGTTCGTGGCTCTCATTGGGAACGTCCTGGGTGAGTCTCCTCCCGGG CAGCCCTCCTAGGGGCTATCACCCCCTCTCCGCCCCGGGCTGAGAAGGCTCTAAAGAGAC CCCTCCCTCCCCGGGAAGCAAACAAAGAGGTCGCTGCTCTCGGATGGGGTTTTCTAATA AAATAATAATAATAGAAAGTTTTCTGATTTTCCGAACCGGGACCGAGCCCTGGAAAG GTTATTCCCTGTTTTGCAGGAATAACGGGGAAACCGCGTTTCTTTTCGAGCACCTAGAT TACAAGCGCAGGGAGAGGGGCCGCGGCAGGGATCTCCAGGTGGATTTTGTTGAGTGTGTG TGTGTGTGGGTGGGTGGGGGGGGGTCAGTCATCCCTTTGTGTAACGTGGCTGGGTGTT TCAGGGGGGTTGGGACGAGACAGAGCTTGCAGAATACAAAGCTACATCCCTAAGGAGCAA GCTCTCTGTGGCTGTGGAAGTCACAAAGCATTTGTGAGCTAGGTGGCATTGCCCTTTGGC GAGGAGGTTTAGTCTCCAGTCAAGAGGTGGTAATGAACCAGCAGGGAGTGGAGACGGAGG CAAAGCAGGGAAGTGCACTCACTCATAGAAGCTGAATTAAACAGGATCCATGCCTGGAGC ATTCTTACATCCATTCAGCCAAATATTTTTTTTTTTCAGTCTGCTTGTTGCCAGGCTCAG GACTAAGCTTAATGCTAGGCTATTTGTCCCGGTCTAGGTCTGTATGCAAACACGGGTTTC CTCGACCCCTCATCCCCCTCCCCCTAAACAATTTCTGAGGGTTGGGGAGGGGGTGAGATG GCAACATGGTGAGTGCGATGATGGAATGTATTAGGGCAGTTGGGGAATATACCTCCAGAA AAGGGGCTTTGGAAGGGAGGGTAACTTGAAATAAATTGTGAATGGAAGGAGAGTGTACC TTGATGAATGAAGAGTAGAAGGCTGGGAGACTTTTCACATGCAGAGGGCAGTGTGGAGGA AGTCTCTGCTGAAAATGACAGGAGATGGAGGAGGCTAGGAGTTGCTCTTGATTTTCATTT ATAAAAGAAGAAGAAGGTGAGTGAGGTGAGATAGGCTGGGAGGCTTTGCAGTCAAAAGCA AAGAACTTGTAGCTGCAATGGGGACTGACAAGGAAATTATCAGGCTTTCAGACTAACCTG ATTTTTGCCTTCTCCCAAGTGTGTTGGTCTGGGTAGAAATCATCCCGAGTAGTCTCTC ACCAACTCAGCAGGCAGAATAGATGATAGTATGTGAATGACAGGAGTTCTCCAGAGTGTT GGTAGAATGTTATTTGAGGAGACAAGAAACCTCTGAGAACTTTAGTACATTTTTAAATAT TATTTTTAGACTGTTTTCCTTTGGTTGATTTAAAAGTAAAAATAAAGGAAATCTTTTTTGG GATACTAACAAAATGAAACAAAAGTGGAAATACACAAGATTAGGATTCTTGTTATAAGCA

FIG. 1H

TAATTCTGTTGATAATAATCCTAATCTTGCTTTCCTTCTTGTTACCCATCCTTAGGA TTACATCTCTTAAGACACATGGCTACCAGCATAGCAACATTTTACTGCATTATGCCAACA CTTATTGATAAGTGAATAATCAAAATTGAACATATATTGAGTACCTACTGTGCCCAGAG CCCTTCATGTACATTCTCTCCCTTAAATATCAAAATAACCCACATTAGCCAGAAGAAGAA ACAAGACTTAGAGAAATAAAATGACGTATTAAGGGACATAATTTAAATTCAGTTCCATTT TTTCTGACCTCAGATCCAGAATTCTCCATTGTTATTCCACTCTAGAGCTAAAAAGCATAT AGAGAATAGATTCTCTGCTCCTGATTGTCTGCAAGTTTATTAGATGTGTTCCTGTTCTCC TCTGCATCAACGCCCACTGCCAATAAAGTACAATGAGGGATTAATGGCACTGTCATTCTC TTCACCAAAAACCTTTCCAGAGAAGCAGTAATTTTTTTTATGAATAGCTATCAATAGTAAC TATTTGCCTTCCTTATTTTAATTTTCGGCTGAATCTTTGTGGTAAAATGTGCTCTTCTTT GTTGTTATTGCATTTTACCTTGCATAGACCTTGTAGTGAATAGTCTCCATATCCTAATT GCATAGTTTAGGGATACATGTTTGCTAGCCTGGGGAGTTTTAGTTTCAAGAAGGAAACAC CTCTACAGTAAGGCTACTTGTTTCATAATGTCAAGGAAGATAGCACTGTCCACAGCCCCA CTTAAAGAAACTGACAGCTATTTTCCTCAGGACTGAATAACACTGAAATCCTCTCTGGTT GAACTGAAATGCATTCTTTTCTGACATACTGCCTGAAAGTTGATGAGGTTTAGGTTTGAC ATTTAAACAAACGAGTAGTGTCGTTACTCACAGACAACTTCCTGCTCTTTGATGTCACTG TCAAATTTGCAAAATGAATTAGATTGAGAATTGCTTCTTTGCCCCTCTGGTATAAGTAAT TTTGCACATAGAGTGGTAGGACAGGATGTCACATGATTTATGCAAAATAAAGATGCAATA TTAAGTATGAAGGTAAAATACCACAGTGTAGGCAGCAGATGTAATCACTGAGCCTTCAGG TCCAGTCACCATTTGTACTTTCATATAACTGCTTGGAAAATCTCAACCTTTTTGGGCTTA CAAATATAATGCCATCAGTTAGAAGTCATCTTCTCCACAATGTCCTTTCATGAAGTGATG TAATAGGATATGCTGTGGGTAGCATAACAAAGTCTTGATTGTCCTCATCTCTTTTTCTTC TCCCCATAGTCCCTCTTTATCACTATGCCACCTCTCCACTCTCATATACTCCTCCCAAAG ATGGAAAGCAGTTTCCTGGGGGGAGTAAAGTTTTAAATAGAATGTTATGAGTATTTACATT GTAAGGTTCTTTTTGGAAGGAGTATCTTTTCAGTATCTTCAGAATAATGCCACCTATAAC CTATTCCTAACTATGTCTTCTACTACAGCTAAGTAGATGTATCAACTTATTCAATTGGTA TATTGTGAGCATTATCATTTTTTAAATTAGTGTGTATATCAGGGGAGCCTCTGGGGAAA TGTAAAGAAATGTGACTGATGTTAATTTTTACTCCTGATTCCTTGAATGACAATTGTAGG GAGAAATGTGTTCTAGTCAGTTTAAACATTAAGTACCTAGGGAAAATGATCAATTTTCTG $\tt CTTCTCATATCTGCATTCAAAGATATCATATGTTTCATCTGGTATGCTTCTGTCATATCT$ GTTGTTGTCTCCATATGGAAAATAGGAAAACATCAGTCTAGCTATGCTTCTTG TGTGCCATTAGCAAGTTATTGAACTATCCAAGTCAATTTTTTTATAATTACAAATTAAAG ATCGATAATGACTGCATTATAGAAATAGTATCAGGATATAATGTACGTATACCCTCTATA AAGACATATAAAGGGACACAGGCATATACATATTTTTCTTGACACATAGACACTAATTAA TGTCAATTTTTATCCCTTAATTTTCATGACTGAACTTTTTGTGATGTGGTGTATAGCCAG CTTCTGCCTTCATGGGCCAGTCTGTATCTCTGTAGCTCTTTATGGCCTCTGCCCCAGCCT TTTCCTTAATTGCATATTTTCCTAAAAGGTGTGAATAAAATGGTGTTGGCACACATTACT CTCCTTTTCCACACTAGCTCCACCCACCCATCTCCTTCATACTGATTGCTTAACATTGCC TTAAATTTGCCTGTGCTGGGTCCCATTCATTTAGAGTTTTTGAGTTGTTAATAGGTTGTT TAAGTAATTTACCTAATAACAGTTTACCCAAGTTAGGTGTGTGGGAATGGGGAAATATTTG TAATAAGTTTGCTTCCTACAGAGTTAGTCTTGTGTCAGATATGTAAGTGGTAGAATTGCA AGTTCATGTTACTCCTAAGCCTAGAGACATTTATTTTCTGCTTCTCCGAATGCCCATTTT AGTTTCATGGGTGTTTGTAAACCCATCCTTACCTACACAGGAAGCAAAAAGGGGTTATTT CTAAACCCTTTTTAGATATAGAAATAATACATCACTCATCTCGGCCAAGACTCAATAGAA TCATGAATAGTGACTGTAAAAGGTAATATTAACTATTAGGCTTTAAACCTATTGTGCATT TTAGTTTTAAAATGCAAACATGCTAATCTGAATAAGAATTAATCTGATGCCTCTACATTT ATCCCAATGATAACTTTTAAGATGGCTATTTCATAGATAACAGCAACATTTATCATGGAC AGACAATAATGAGAATAACATGTGCAACTGATAATTTAAATGCAATGAGTTATTTCTGTA TTTGAAAAATATTTTGGGAAATGGGATAATTAAAAAATACCAGTTTTCAAGAGACCAA ATCTAAAACTCAAACATAAACACAATGCTCCAGTTTTTAGAAAACTGTCTTGATTGTAGT AGTGCCTACATACTAAATTGTATCATATGATTTATATTAATTTTCCTTATTTTGTATTTT

FIG. 11

AGATTATATTTGAAAATTTTCATGTACTGCAGCTATGTTAGCATCTCAAAGTCTCCATAT TCTCACTCCGCTCCGAAACATCCACTGCTGATGTTATTTAACTAGTGAAAGAAGATCCTT CCATGTTTCTTATAGCATTCTGACATCTTCTCCACCCTAAGGAATGCTGGCTTTATT AAGTATGTTTCAGTCAATGACATGTGATTGGTGAAGCTGACGGTATTTGTCTTCAGTTCC TTTTTTCCCTGCAAAGGAAATTTGTTGAATATTTATTGGGTACTATATGCCAGGTACTAT ATGTCAGGCTCCACTTACATATACTCTATTGATGCCTTACAACAAACTTATAATGAGAAG ATTAATAGGTTTTACAAATAAGAAAAATGAATTCAAAGAGCAATGCTAACTTACTCAAAA GTTTAGTCAGGCAGTAAATAGCAGCACTAGGTTTCAAATATGGATTTAACAAATTCCATG GTCCATGCTTATTCCATTACTTCATCCTGCCTCTTTCCTTAGCTTCTAACCCTGACTGGA GATGCATAGGCAAAAAGAGGAAGGAAGGAAGATACTTAGATGTGCCCTCTAGACAATTTACA GAGTTGTTTGGGCATGTTGCCATGCTGTTTTTCTGATAGACTACAGTTCTTCAGCTCTGA GGATGAGCTCATTTGATAAGCCAATCAAGGTCGGGCTAGGGTTACTTTACAAGAGAAAAT TTCAAGGTAAAATAGGTGCTGCCAAAAATGCTTTTACCTGTTCAGGGGGTTGACTCACTG GAAAAAAATGTTAGATAATTGTGGCCAAGGATTATTTTGTTATTGAAAGTGCTATTTTT AGACACAATTTGAGCCTGAGAGCCTAAACACTTAACACTTCACATAATCTACAGATATTT CTTGCTCTGTCACCCAGACTGGGGTGCAGTGGCACAATCTCGTCTCACTGCTGCCTCCAC CTCCTGGCTTCAAGCTATTTTCCTGCCTCAGCCTCCCCAGTAGCTGGGATTACAGGCACA ATCTTGGCTAGGCTGGTCTTGAACTCTTGCCCTTGTTATCTACCCACCTCAGCCTCCCAA AATGCTGGGATTGCAGGCATGAGCCACTGTGCCTGACGTGAACAGGTCAATTTCTATATC ACATCAGACATGAAATGACCTTTAGATACTGACTTTGAAAGAGTTTGAGATGCTATTGGA TGAAACACATGACCCATATGACCAGTCTTTTGAATTGCTGACTCTGAGTATAAAATGTTT TCATTTCACCTTTGTTCACAATGAGAAGTGATCTCTTAACCAAGTAAATGAATTAAATCG ATATTTAAAATAACATTAAATTTCTTGCCAGAAAAACTGTTCTTTCATAAACAAAAAACA AATTGCTCAAAATAAATGACTATATCTTTATTTCTAAAAAATGTTTAGAGATTATTATTA TTGGGTCTTTACAAGTAATTTGCCTTCAATACTAAACACATGAGAACAATGTTTAATATT AACAACATTATGAGGTAGGTCTTTTTTAATGAAAAAAAACTCAAGTGCTTGAAGTGATTT AAAATCACTGTGGAAGAAAAGCATGGGCATACAGAAAAGCCAAGTGGTTGTGTCTAGCT TGGGAAAAGCTTGCAAATTTCCTGTATTTCAAGAGGCCAGGATGAGGTGTGTAATTATCT TTTACTGGTCTTCAGCTATCCTGTCTTTGATATGTGATTGTGTCAAAACTATGAGGAAAA ACTCACATTAACAAACTTCATAAACTTGTTAAACATAAAATAATTATCGATGTTTTAA TTTACAGTAAGAGTTTATTCTTACAAGTCCTTAAATACCCAAAGTTCTTTCAGTTATCAT AGTCTTTTTCAGTAGACAGAAATCCATGTGGACTGTTATTGTTCTGAATAGCTAGGCTAT GCCATAGTAGCAAACAAACCCTGAATTTTCATTGGCTTAGTATCACGAAAGTTTATTTCT TGCTCATTTAACATCTGAGGTGGGTTGGAGAGTCTCCTTCATCCAATGACTCACAGTTCA GGCAGCCTCCACATTTTGTGCACTATCCCTAAAAGGTGGACTCTGTGGTAATCAGTTTCC AATATGGCTTCCAATGACCGCCCCCGGGCCCCGGCCCCACTTCCTGATAGTCACATCATC GTGTAGTCCCTTTGCATATTATGCCAGAATTGGTCTGGGTGACCAACAGCTCATAGCAGC AGTGAAACGATGTCACTTTCAAGATTACATAACAGGAGCTTACAGCTTCTGGCTCAAGTA CCCACTTTCTCTCTAGCTCTTGGATCTCTTCTTCTGGAGGAAGTAAGCTGCCTTGTGGTG AGCAGCTGTTGGCTGGAGTTAAAATCTCCAGCCAGCAGCCAGAGAGAAAATACGGTCTGT TAACAACCTCATGTGTGAGCTTGGAAGCAAATCCTTCAGACCAGGTTGAGTCTTGAGGTG ACTACAACAGCCACTACCCCAACCCACCCCCAGCTTCAGTGCAACTTAGTAACAGACACT GAGTCAGAACTATTCAGCTAAGCTTCTTGCAGATTCCTGACCATTCAGAAGCTATGTCAT AATAAATTTTTGTTGTTTGACTTCAGTTTCGGGATAAGTTGTTGCACAGCCTCTAAAGTT GTGAACTAGAAGAAGTATACTGGCTCTTAACCACCTTTGCCAAAAATTAACACTTGTCAG TCATGGTCATATTCATTTGGTCCAAATCAATCATATCGTATCAACCTAACTACAAAGGGG ATTGGGAGATGGTGATGTCTCTGTCACAGAATCTATATAATAGTTAAAAGTATTTTTAAC TTGCATAGACTCAGAACAAGATAATTTGGAGGAATTCAATGCTTAATGGCATACCACTAA GATAAGCTGATAGATATATCGTTGCGATTTGGGTCTCTGACAATAGAGGCAATTGATAAT ATTAAGAGACTATGTGCCAATTATTGTGCTTGGATTGAGGGTACAAAGGTAATAGAATCC AAGGAACCTGCACTCTTTTTGAAAGATAGACACATAAACACATACTTTTAAAATAACGTG GTAAGTGCTACTATGACAGATGGTTGCACAGAATGTAGTGGAAGTATTTGAGAAGGACAC

FIG. 1J

TTAGCTCTGCTGGGGGATTAGAGAGAGATACAGGAGGAGATGACACCTAAACTGAGTTTT AATAGATGAATTCAAGTTACCCAGGTGAAGAAAATTGGGTAAGGATGTTCTAAGCAGAGG AAACAACATAAGCAAAATCAAAGAGGGGTGAAATAGAATGAGCTATGAAGAAAGTGTTAG GCAATTGGGTAAGTCCAATGTAAGTGCAGATGAGGAGAGTCTGGAAATGAGGCTGAAGCA GTAAATAAGGATTGGCCATAAAAGACCTTGTGTACAATTCTTAAGATCTAGGCTTTGACA CTGTTGTTTAGGGGGGGGCTGTTAAAGGATTTTAAATTAGAGTACCATCATTGGTTTGCAT GAAGGTTTTCACACTGGGGTTTGCATCCTGTTTTGGCAATAAGCTTGTTTTAATGAAAAC AAACAAACAAACTGACAATAAAGAACATAATCCAAATTCTCCAGATAATTACTTCCAGGA GGCTTTCTACGTGCTGCATACAAAACAAAGAAAGAAAAACATAAAGTGAGAAAACGAAGG AAAAACAAGGAAAGAAGAAAGAAAGAATACATATTGGAAAAACTGTTGCTGTTTTTGT GATAAAGTCTCACTCTGTTGCCCAGGCTGGAGTGCAGTGGCGCCATTTCAGCTCACTGCA ACCTCCGCCTTCCAGGTCCCAGTGATTCTCCTGCCTCAGCCTCCCCAGTAGCTGGGACTT ${\tt CAGACATGCACCATCACGAGCAGCTAATTTTTTGAATTCTTAGTAGAGATGGGATTTCAC}$ CGTGTTGCTCAGACTGATCTTTAACTCCTGAGCACAGGCAATCCGCCCACCTTGGCCTCC CAAAGTGCTAGGATTACAGGCGAGAGCCACTGCACCCAGGCGCAGGTTTTCTTTATGATG TTTTAATTATCTTTCTTGGAACATATATGTATGAATCTTGCATGCCATAGGTCTATTA ATATTTTCCAATATTCTACATGGTTTTTTACTAAAATCATTTTTATGATTAGTTACTGAC TGAGGTTTCAATGCATCACTGTACTCCTAGCTATCTCTCATTTTAGCTTTTACATCACAT TTTGGCCTCACACTGAAACACAAAATATTAAAAAATTTGAGATCTAATAAACAATTTTCAC ATTTTCCAACTAAATCCCCACTTCTTTCTAAATTTTCTACAACTTTCTAAACATTCTCAC GACCCCAAGTGAGCCCTTAGGGAATTTCCGTGAATATTTCCCTACAGGTTGGCATGGTAA CACACTTCACAATTTCTAAATCTGTGGATAGTTTAGAAGCTTTTATTTGCTGTTCCTAGT TCACAATGGAAATACAACAATGATTAAAAATTATAATATCCTTTTGTAGATTCTTAGCTT TTATTCCTACTCAGTGACTCTAAAATGAATTTATAAGGCCCATGGTTTATAACCATGTGA GGCCTTGATTTTGTCACTACATTGCTAGAAATGGGGTCAGAAGGCCACCAGCTTTAATAA TTTAATTCATCAATTCGGAATGAATTTGATGAGTCAACCACTTTGGTAGAGAACCATATT GCTCATAAATACTGTTTTGAAGGCAATTCGTCTTTCATAAAATGTGAAGATTGTGCTGAT CTTTCTGGGCAGGGTTATGGAGGTGTGATTAAATGCTTAAGAAACCATTTTGTTATTATA TTAAACCGAATCAACTTTTTATTATTAAAAAATAGATAAAACTTAGCATCCTCAATTATA ATACTTTATACAAAAGTTTCCCAATTTTATATAGACTGAAGATAAAAATACATTAACAAA TCTTACCAGCTGGTTCAGGAAAATAACTTCATAATTATTGAGACATTTATGTGTTTTGGGC TTGATTTATACTTTGGACACAGGAAAACCTAGAGAGATCTGGTTCTTTGAAATCATCAGA GATGGTGATGGTGACTCAGAGATTCCTGAAAATCAGTAAGATTACCCTAGTTTATAGACG TATGTGTTATTTTTCCCCCAGGCATAATGAACTTTATAACTTGTCATTGACAAGAAGCC AAACATGTACATACCTCACACATGTGTACACACACAGTTTGGGGATTGGATGATATGAAT AATATAATTAATACACCCTAATTTTTCATGCAGGATTAAGAAAGTATCTTCCAAACATTA AAAATGCTGAAAACTGGACATAAGGCCTTGAGTTTCCCAAATTCAGGACATATTTTCAAC TATCCCCTGAGTAAATGAACTATAACATTTACAGAAGTAAAAATGATAAAATACACTAAAG CTGTAGCATGATTCCTTGAATAGACAATATTCCTTGACAATCTTTCTGTAAACA GAATACAATGTTTCCCTAAGCAATATATGCGTGCTCTAGAGTTTTCACAATTTCTGATCC TCCTATGACTGGCTCCTGCTCAGCTCACACTGCACTTTCATGGAAGTTCTCTTAGAATGC CAGCTTTGAATCACTGCTCCCTCATGTGCTGTGTGTGATAGCATCCCATTTTAGTTTTGT CATAGAATTGATTACCATTTCAAATTGAATTGTTAATTTATTGTTCATTTTTCTGTTGTC AAGTGTAAGACTTCCAGCAGGAGGAATTTTTTACATATAAGTACATTTTTTAAATTAAGC ATTGCAGGCTTTAAATTTCTTCTATATAAATATTTAAAATAAAGCTTCAATAATTTGAAT TGCTTTTGTGATTATTTTGTTTTATACCTTGAGTAACTTATACATCAACTATTTTGTAGT TATTCTAGTAATGATTATGAAAGACCATTTGAAAATCTTTCCCCAGCACTGAGATCTCCT TGACATGACTAAGTGATTTATACTATGCAATTATATTGCTCTTCTCAAGAAAAGCAAAAT GAAATTTACAAATTTGGTAGCTTTTTGTTCTTTTGTTTTCTCAAGTAAGATACACCAAGA TTTCTTTAAATGATACGCTATATTTCTGCAATAACTGAGAAGAACATGTAATGTGCAAAA

FIG. 1K

CTCTTAAACTCTTTTTGTTTCAAAATAATTCTTGGTTGTTTTTATAAAAGTCTAAGCAAA TACTTAATGAACTGTGTCCCAAATGAGGTGAAACAGCTGTGACAGAATGTTACTATGACT CTGTACTTTCTATAATAAAAAGGGACAGACATATCCTCACCTGAGCCTTGGGATGTTTCA GGCATGCCCATAGAGCCTAAGCTTTAGGAATCCTCTGTCATTCTTTTCCATTGCCAGTGA CTTGTGCCAATTCTAGGGTTCTGGACTGTGCAAACAATGGAAAAAATAATAACACTTTCA AGAATAGGAGAAACACTAATCCCATCTAATTCTGCCTTCAAACTCCTAAAATATTCATCA CAGGTCTGGTGGCTCATGCCTGTAATCCCAGCACTTTGGGATGCTGAGGCGGGTGGATCA TGAGTTCAAGAGATCGAGACCATCCTGGCCAACATGGTAAAAACCCCATCTCTACTAAAAA ACAAACAACAAAAAATTAGCTGGGCTTGGTGGCATGCGCCTGTAGTCCCAGCTACTTGG GAAGCTGAGGCAGGAGATCACTTCAACCCGGGAGACGGAGGTTGCAGTGAGCCAAGATG AGCAGATCCTGGAACAACTGAACCAAATTTATTAATATGTATTATTACTGAAAATCAGTA ATGAACAAAATTTACAGAATGGGCTTCTTGGAGTTGTTACATTTCCCTTATTACATAACT $\tt CTTCAATAAAAGTGTTTGTCATACCTATTTTAGTTAATTCTACAACAACTAGTGTGATAG$ GGCTATTATTTGATCTTTTTTTTTTTTTTTTTTTTTACAGGTAGTGACATTCAGTATTA GACAGCTGCTATTGTGTTAGTTGTCTGAATACCTTTACATATTATCAACTGGCCTTTTCA TTCCTGAGTTGTGAGTAAATGCTCTGTCTCCCAGACTGGAGTGCAGTGGCGCAATCTCGC CTCAGTGCAAGCTCCGGCCTCCCGGGTTCACACCATTCTCCAGCCTCAGCCTCCCGAGTAG GAGACAGGTTTTCACCATGTTAGCCAGGATGGTCTTGATCTCCTGACCTCGTGATCCACC CGCCTCAGCCTCCCAAAGTGCTGGGATTACAGGCATGAGCCACCACACCCGGCCATAAAT GCAGTCTTGTGTTCCCCACTTCCATTCCTCCTTTGACAGTACAGCTATGCTAGTCTGCGT AGCAAATTGAAAAATATGACCTGTGGGATTTAAACAAAACACAGTGTCATACACATTTT CTGGTAAACTTAACCAAAAGGGACTTGGGTTCCATAACTAATCACCAATGCCTCAGTGAT CTGTAACTCCTTGTAGGTACCTGATCACAGTTACTAAAGGGAAAGAGGAGCGAGGAATAC AAGAGCAAAGTCAAGCCAGACATAGATTTTATCTCTTTGTAAACAGGAGTTCAGAAGACC GCTCTGAATGCTGAGTTAGCATCAGCAATAATAGAAATATATGCAGATTGTTGATTTGAA GTCATGCAAAGATATCTTTTTCATCCAAATGGAGGCAAAAGCATCATAGAGCACCAGAGG GCTAAATCCAACTGTAGCAGCAAAAGGTACACAGAAAAATAAAGCATCCTGAACCAACGC ATATTTATTGAACACCCACAATGTGTAATCTGTTCTATTACATTCTGTGGAGGAAATACA GAAGTGAATGAGGCATGGTTCTTACCTACAAGGAATTTCTAATCTTGTGGGGGAGACTAA CATGTAAACAATAAACTATAGTATGAGGATTACTGAAGAGGCATATGCTAAGTCTCAGAA TAGCATTTTGAGTTATTGTTGACATGTGAATACGATTTTGAAAAGTTCCAAAGAATGAAA AATTCCACCTACATTGGTGAAGTACTAAGATTAAATGCATGATAGCTTGAAGACACAAAA ATAATTATTATAAACCATTCCAAAAATCATTCAGGGAATTCCAATAATACACAAGTTTT TAAACACATTTCTGGGTAATTTTGAGTAATAAGGTCTTAATCTCCTCTACTGCTTTCAAT TGTTTTTGTGGCCTTCTTTATTTTGTGGGTATCTGGCCCAGTCTTGTCTGTAGTGTATTA TGGTGGATTGAAACATGTTTTGCAATCTCTGGAGTGATTTTAAAATGACTTGTGTT ATATCAGAGTTTCCTAAAGGGAGATTAATTTGGCTTAATGGTAAGAACGGATTAAAGTTA TGAGATACCAGACACTGGGAAAACAGTTAGAAGCCTGTTGAGACTCTTCAGGGCAGTTGT GGCAGTCTGAAGATGGAAGTTGGCAACTCATCAAATGTGAGAAATTTATAGGAACAGAAA AGAACCTGCTGATTAATATAAATTTTCTGCCAAAGAAAGTACAGTGGCTCTCCTCAGCAA ACTAACATGGGAACATAAAACTAAACACTGCATGTTCTCACTTATAAGCAGAGCTGAACA ATGGGAACACATGGACACAGGGAGTGGGACATCACACACTGGGGCCTGTTGTCGGGACTA TGGGAGGGAGACCATCAGGATAAATAGCTAAAGCATGTGGGACTTAATACCTAGGTGATG GGTTGATAGGTGTAGCAAACTATGATGACACACGTTTACCTATGTAACAAACCTGCACGT GTATAGAAGGGTCATGATGAATTGGACAAAGATACGTGGAGTTTGAATTGCTAGAGGAGT ACCCACGTGCAGTTGTCCAGCAGAAATCAGGGCTTGTTCCCCAACATGCTATTCACAATC

FIG. 1L

13/51

ATGTGAAAATGTTTTAGAAAATGTCTTCTGGAATAATTAAAAGCATACAAGGGAATGTAA ATCTCTTAATGTGACAAGACCTTTTTGCCACAATAAACAAATTCATTAGTTCAAAAAAATA TTTATTGTGTGCCTATTGCAGCAAACAAAACAGACGAAGCTCCTTCTTGTAGGGAACTTA TACTCTAGTGATATTTAGTATATTTTTGACAATTGAACCAACAGGATTTGCTGACGGAT AGGTGCCATTTACTGAGATACAGGGCCGTAGAGGAGGAGTGTGTTTGCAGCAGGGAAGGA GAAGACTCAAAATTTGGTTTTGATCATACTAAATTTGATATAGTACAGGTAAGTGTATGG TGGCCATTAGAACATGAAGGTAAGAGTTTAGATAAGGAGACAGGTATGGTGAAATACATC CAATATTTATAACCAATATTATCTTTTGTGTCTGTACCTTTTTATACATTCCCCATATAT ATCAAAGACTATAGAAGGGACTGGATAGTGAATAAGTGATTATACATAAATTCTTTTTTA CAGATTATTTTGCTCTTGATTTCTCCTATGTAAATCATCACAGCTACATTTTTTAAAATC TTAAAAAGGATTACTTTGAACAATGCATTTAAACATCCAGAAAACAAAAACAGGAGTGCA TGGTAAAAATTCTGATTTCAGAACGTATGCCTGACTTATCAAGTCAGAATTTCAGGGAGT GAAGACCCTGGAATCTACACTTTAAATAGAGCCTCAGTTCACCAAGTATGAGAAGTCCTG TAACAGGGAAAAGTAACCTCCTGTTATATTTGATGGAGGCCAATTGACAAGCCAAGTAGT TTTCCATTTGACAAAAATTCTATTGTACCAATGAAGAGCTATCAGAGGGGAGTAGATTAA AACACCTCCCTTGAAATGGAATTTGGCAAGAAAGCAAGAAATTACAGCAAAAAGACCAAT AAGAGGAATTAGGGGCAATGAAGGAAGGAGCAAAGATGTGGGAACCCAAAAAGTTTTCCT TACATTGGTGTAATTTATTATTATTATTAAGCCAACAATATACTTTTAAACTTATACAACT TTGCAAAAAAGTACAAATCAGAAGTCTGGGCTAAGTAGAATGCATAATAGAATCAGTAGT ATTAAAATGAATGTCACTTCTTTTTACCATGTGTCCTTTAAATTATTAAAATCTATACAC ATATTGCTATACATAGTAAATATAGTTAGTCAATTATGTCATGGAAAGAATTGAAGGGTT GTTATAAATTTAAAGGTGTTTCACTATACAAAAACATTGTGAAATACTGGTGCTGATTTA GTTCTAGTATCTCTGATATATTAAATCATAAATGTCAGGAGTTATTGGTCACAAAATAAA CACCAGAATTATATGACAGTCTAAAAACAAAAACAAAAACTTCAGCAACAATATTGAAG ATATGGAAGTGCCAGAAGAATAAGGATTAAGACAATGAATAAAAATCTCTTCCAAGGACT GGTCTACACTAAGAGTTTAGAAATGCATTTTTTTTTCACAGAAATATCCTTAATCCTCTA TATAGAAATGAGAAAAACATAAGACTTTAGCAAGCTCCATCTAATCCATTTGCAGACA TATGGTTACCTATCTTTCTTCAATATATTGGAGTTTGCAAATATTCTACCTTCAAAGAA TAGGTGTTACCAAAACATTGTCTGCAAGATTTCTAAGATTTGAAATATATTTGCTATAGT AGGTTAGAGATGAGACATTTTTACTTTAAATTGCAATAATTCAGACTTAAAATATAAAAT TGTGTATACTCATATGAACTTTAAGGAAATATCAGAGGCATCAGTAATAGATAACTTGCA TCTCTTTTACATTCAGTTCAAGCTACTCAAATTTTAATCTTTTGTTTTCATTCCAACAAA AAAAATTAGGATCTGCCTTGGCTTTTGCTAAGAAAGTAATTATTGGCTGGACATGGTGGC TCACATCTGTAATCCCAGTACTTTGGGAAGCTGAGGTGGACAGATTGCTTGAGCTCAGGA CGCGCGCACACACACACACACACACACACACACAAATTAGCTGGGAATGATTACACGC $\tt CTGTGGTCCCAGATACTTGGGAGGCTGAGGTGGGAAAATCACCTGAGCCCAGGAAGTCGA$ TTATCTTCAACACTGTGCATACACACTTTTCTGCATCTAGATCCCAAATTTTTGTTTTGT ATTTACATAGAACATTGATAAGTAAGGTAAGTATTAATTGATAAAACATTTCAAACTCAT TTTTCACTAAATCCAATGGCCTTCCTCTTTTGCATGAAGTCTCTAAGAATCATGTTAATC TACATACTCAATCTACGTAACAACTGGATATATCCTGTAGTTGTTGCCCATTTTTCTGCT **AAATGTTATCTTTAGCACTAAGCATGAGTATGAGGAAACAGTATCTGTGCTCAGATTCCA** GAAATGAAGAAAATGTACTGGAGGTCTTTTGGATAATGGCTACAAGGTCACAGGGACTGA CTCCTTTGGAAGCTCAGCGATAACCATTTTCAGAGAGAATATGTCAACATCTTTCAGTCT AGAACTTGATGTTCTGCTGAGATCTAATCTGGGGGTGTCCTACTATTGAATAGGTATAAA ATATTGAAAGTCTATTAATTGGCAAGCACTCTTCTGACATTAGAAGGAGCAAAGATAAAA AAGATATTATCATTAACCTCAAGGACATGACAGCATCATGGGAAGGCCAGAAATGCAATA TGTTAAAGTAAAACACAGTGTAGTGTTTACTACTAAAGAGATATAAACAGAGTACTGTGG TCTAAAATCATATATATAACATTTGCTTAATGGATGAGAAGGAAACTTTAACTTCAGGAG GCAGAGCATTAAGAAAGTGAATGACAGGAGGGTCAAAAGAAAAAGCCGACAGTGTTGCAG AGGCAGGGCATAAAGGAGCTAAACCTTTGCTACCTTCAGTTTTTATTATCCACAGAACGA

FIG. 1M

AATTTATGGGGGCATTCCTATGGTCCTCACCTCACCCCATTTTTCTGTTTTACCTATGAA ACTTGATCAAAATACTGTCTCCACATTTCTCATAAAATACATTAGTTTAATTTTCTACTA TTACTTTCTTTTAGTTGATTTAAAAAAAGGTCATTTATGACCTATTTAGGTTAGCATCAT TAATTTTATCAATGTAAGAATATGGTAGTACAGTGTGAATTCCATTAATGGATATGTTGA TACCATGGGTTTCTCTGACCTTTCCTCTTCCGCTCCTCCTGATGATTGGTTCTGAGCTT TTTAACACTGAATAAAGATTTTTTTTTTCTCTAACAGACTTAAAAATAGTGCCCTAAAAAT AAAAGAAAAACAAAAAACAAAAATAAACATTGTGTCCTACATTTGTATTAACTTTCTTA TTGAGTTTTATTGATAATACCTTTAGGATGCATGTATTATTAGAAACATCAGTTATTTAC AAGTTCACCTATTTAAAAGTCTAATAGGAAAAAATATTTCATGTTGCTAAGTATGTGACT TCCCTTTAAAAGATAATAATGCTTTCCCTTTAAACAACAATAGTAAAAGAAGTAGAGTTC CTTTTAAACACATACTTTTATATTATAACCCATTCTGTTTAAAAAATAGCAGGCATATAA TCTAGAAATGCAAATAATTTAGTGAAATTTTTTAAAATTATTCTACATATAATTAAATATG GATATTCGTTTTCAAATATCAAATAATAAAATATGTCTGAGATGCTGACTAATCCTTAAT TATAGGTGTGATTTCTACTTCACCATCAATACTATGGTACTCCAAATCTTAACATGAGTC TGATTTTCTAATAAACATGATGAAAAAAGTTATGGAAAAATTTTGAGATTTACTTTGGGA GGTTCTATTGTGTTCTGTTCAGCTTCATAATATTCAGTTTCTATGAGTTTGGTATTTAAT TATGTGTGTTTGTCATTCAGTAGGCTGGAAGTATGACCATTGGGAAGATCAAAACGATAAG ACATTAATGACAGTGCTTTATCACTGAATCTAGTACTTTTTTTAATGAAAGAGATGTTGG CCTCTTGTATTGTTATAAAACAACACAATTTTATGGCTTTAAATTAAAGTACAATCATAA CAGAAGACAAAATTAGATTAAAAAACAAACATGGAGTGACTCATATAAAATATTTAGAAA CCAATAATACAGATAGAGACACATTAGTTCCTCTAGACATTGTGTTTTCCAGTAAAATGA TCACCAAACTTACCAGGAAAATGATAATTATCAGATTATTTACTTTCAGAATTAAAGGCA GGAAGAAAAAATGAATGAAGAGGAAACACAGTAACCATATAGGACAATAAGAGTGAA TGAAGATAAAATGAAAAATCAATAAGATATCGACTTTCTTAAAAGACAAATATCACAATA GGAAACACCTCAGAAAGGGAAATCTCAAGAAAATAATAAACTGAAAGAAGAAAACATATC AAAACAACTTGAGGACTGACAAAGTTTTAAAATGTATTTAGATAAAGATACCATGAGGAA AGTGATCAAGGTGTTCTAGGTAATCACTGAAGATAAAACTAAAAATAGCTTAAATTAAAA TGAACATCTTAACAATGTCTCAAATGTCAGGAATTGATCCAGTTTTTTGGCTGCACAACAG AGTGGCTATAGTTAACAATAATTCACTGTATATTTCAAAATAACTCAAAGAGTAGAATCG GAATGTTGCTAACACAAAGAAATGATAAATTCTTGAGGAAATGGATATCCCAATTACCCT GATTTGATCTTTACACATTGTATGCTTATATAAAAACAGTATTCATGGCCGGGCGTGGTG GCTCACACCTGTAATCCCTGCACTTTGGGAGGTCGAGGTGGGCGGATCACAAGATCAGGA GATTGAGACCATCCTGTGAATGGTGAAACCCCGTCTCTACTAAAAATACAAAAATTAGC CGGGTGTGGTGGGCGCCTGTAGTCCCAGCTACTGGGGAGGCTGAGGTGGGAGAATGG CATGAACCCAGGAGGCAGAGCTTGCTTGCAGTGAGCTGTGATTGCACCACTGCACTCCAG CCTGGGCGACAGAGCGAGACTTCGTCTCAATAAAACAACAACAACAACAACAACAAAAAAAC ATTTAAATTTTTCCTATAGCATAGAGATCTGTAATTAATACTTGTCGATCATTGTTGTTT CTGTCTTCCCAACAACTACACTCCTGTTTCTTCACATTCCCCCTTCTTCTAACAGCACTA CATCTTTCTTTAGGAAACTATCCTTTTGCCATTTCATGTATATGGTGGGGTGGGGGGAGTT ATCAATCACAGTACCCCAGCAGATGGGACCAGAGGCAAAAATGCCTGACCTTCTCCCATC CCCCAACCACAGCAGCAAATGAATTATAATTTGATGCACAAGGAAGTATCGGAGCTTTTG TGTTGGGTTTTACATATCACCTGTGGGAGATAAATGAACTTTTCCCCACCTAACCTTTAG CCACTTGGGATGATTAGACATAGAGGTGCCTAAGATCTTTCCCTTTGCCACATTAAAAAC AAATCATCTATGGCACGAGCATACAAGACCAGCTTTCAGAGACACAAAATGATGGAGAGA ACCATGATACTAGTTTTAGACCTAGTCACTGAGACTTTCTCTGCTCCTTCCCAGTTACCT GAGCTTTATTTGTTTACATTTATCAGATTTGAATGGCTGTACTTCAAAGTACTGATTAA AATAGGAACCAACCTATATGATTCAGGTGGTGAGAAGAAAAAGAGAGAAAATGAGG TTAACAAAAGAGAATAAAGAAAAAGAAAGAAAGAAACAAGAAACTCTGACTACCTCTCC TCTTTGACATAGTTTACACTTCTGACAGATTGTTCTTCTCTAAATTTATGTAGAGATTAG AGTGAGGATGATGTATGCACTGTAGCATGGGTGGTCTTCCAGGAAGCCTTGACTGAATGA

FIG. 1N

GGCAAGGAGTATGTTGCTCCCTCAGTAACCTCAAATTTACCTGCAAGCCTGATAAAAATC TAACACTAACACTAAACCCAATCTTATCTACAGCCCTAACTGCACCCTAATATTAACAAC CCTACCTCTGTACTTCAAAACTAAACTAATTCTGATTTTACTCCCATCTGCCCCTTTTA CCCTAAAACCAACTGTAAAACTAAATTTAACTCTAAACGTAATCCTAAAACTAAGAATTA ACTAACAATTTTATCTCTATACCCAACTGTTAACCCCAAGCCTAACTCTAATCCTATCTC TAACCTAACATTAACCACAAACCTACTTCTAACTCTAATCCTAACCATAACCTCAAATCT **AACTCTGATTCCAATTGTAATCTAAACACCAACCCACCCCTACCCTTTTATCCCAAATCC ATCTGAAACCCTCATCAGAACACAAATTCCAATTCTACTTCCCACCCTGACTCTGACTCT** AAACATAGGCCCAAATATAACTCTAACTCGAAGTCAAAAACTTAACAAACCTTATCTTGA AACTCAACCTTTACACTAACCCCAATTCTATCTGTAATCCTAACCCTAATATTATCATCA AACCTATGTCTAACACGACCTCCAACCCAAAACCAAAACTAACCTCAGACCTAACTCTAC ATCTAATTATAACCCAAACCCCAGGCTGCTACTTACCATAACCCTGAAACTAAGCTTGAT CCTTTCTCTTTTTTTGAGATGGAGTCTCGCTCTGTCTCCCAGGTTGAAGTGCAGTGGCG TGATCTCGGCTCACTGCAAGCTCTGCCTCTCAGGTTCATGCCATTCTCCTGCCTCAGCCT CCCGAGTAGCTGGGACTACAGGTGCCCGCCACCATGCCTGGCTAATTTGTTGTATTTTTG GCAGAGATGGGGTTTCACCCTGTTAGCAAGGATGGTCTCAATCCCCTGACCTTGTGATCT GCCTGCCTCGGCCTCCCAAAGTGCTGGGATTACAGGCATGAGCCACCACGCCCAGCCGAC CCTTTCTCTTAACCTACACTAACACTAACTGTAAACCTAGCTGTAACTCTAATTGTAAAC CTAACCTGATTACTCACTACAAAGGTCCCTCTAATTCTAACAAGAAACTCAATCCTATCT CAATTCCACCCAACCCCAAAAGTAAATCTAAACTTAAACATAACTCAAAATGTATCTCAA ACCTTAACCTTCACGAAACTACATGTAGCACTAATGTAACCCTAAGCCCAATCCTATCAG TAACACTAATGCTAAAACAAACCCCAATCTGTATCTCTACCCCATCACTGACACTACCCA ATATCTCCAACCCTAACCCTAACATTAACCCCAAACTTATCATTAATCATACATCTAGCT CTAAACCTAACCCCAACTTTAACTCTTACCCTAGCTCTAAAATTAACCCCAACACTATCT CAAACTGTAATCCTAATGCTAACGTTCAGGCTACTTTTAACCCTCACCCTACACAAAATC CTGCACCTAAACTCAACCCTAACTTTAAACCTACCTCTAATCCAAACACTAAATTTAAAT CTGGATCACGACTTGGGCATACTAGCACCCACTAGTGTTCTGGGTGTCATTTCTTTGCTT CACTCTCATCCAGCTTTCTTTACTAATTTTGGATAATGAATCAGAAAATAGTGGTGTGGA ATCAGGGTCTTTGAATTATTGTATTATCCAGAGTTTGTCCTGCTGCAAATGATAAAACAC TTATTGGCTCAAGGAACAGAAGAATTCAGTACTGAGTTTCACAAATACCTGGATTCCCTA TTGGATTCATTGTTAGGTTTCCTGTGGCAAGATGAAATGGCCTCAGGCCTGTAACACAAT AGGATCAATTACAACAGAAGATAGTATTTCTGTTTTCCTGGTTGCTCAAGCCTAAATTCC AAGATTAGTTTATATCAAACCTAGTTAGTTTTGCTCATGTAAGGGATTACTGCAACTGGG TACACTAATATGAAGAGTGGGAGAGTTGGTTAAGGGGGTTCTCTGAAAGGAGAATTAGGT CAAATGTTGAGGAAGAATTTTCCTTTATGATCATCTAGCCCAACTTTTTAATTTCTAATT TTGTGGTTTTGACCAGTTTTTTTTTTTTTTTTAATGCAGCATGTCATAAAGTTGGGAA TACTTCACATTTTGTCTTTGAAAATTTGGAGAGTACTTAAAAAGATTTACAAAGGGGAGG ATTGAATTATTTTAGGAATGTAAATGGTGGTCTTCTGTCTCAGGCAATGTAGATGCTTGC TAGAAAACAGCTGACTCATGACTGTTTTCTTTCTAATTCATTAATATGAATTATTTCAAA CTGCAAAGTTATCTCCTTTCTTCTCCTAATCTATCCACTTAGAGTATACATGTTCAAATT AATGTATTGAACTAATTTTTCTAGTAATACATTCTATGCATTACAAAAATAGCAGTGGGA AGGTGAAAACAAAATGCAGTTATGCATTTATCTCTAAATGTGTTCAACATCTCTTATGCG TGGGGAAAATTTAAAGCTGGCGAAAAAGGCTTCATCATAATTGACAATATGGGAAAATAC TGTATTAAAATCCTAGGTTTCTCCCTTGTTTGCATGAAGGAAATGAAAAATATATAAGGG AAGGATTTAATCAGTCAGGCAAAAATCTAAATTCATCACAGGTTTATTCACTGCATACTA TAAATGAGCAGAGTGTGGTGATTAAAGGTTGGTATTTGGTGCTGGGTAGACCCAGCTTTG CCACTTACTGCCCAAGTAAATATTGCCATCCATCAGATATCTCCACCTATCAGACCCACC CTGTTGTAATAACAAGATTAAAATCTGTATCACTAAAACTTTAAAAGAATTTATAGCCGA TGTGTGTGTTTTTGAAACGGAGTCTTGCTCTGTCGCCTAGGCTGGAGTGCAATGGTGGGA

FIG. 10

PCT/US00/23021

TCTTGGCTCACTGCAACTCCCGGCCTCCTGGGTTCACGTGATTCTCCTGCCTCAGCCTCCT GAGCAACTGGGAGTACAGGCGCACACCACCCAGCTAATTTTTTGTATTTTTAGTA GAGATGGGGTTTCACTATGTTGGCCAGACTGGTGTTTTTTGAAAAGACTTTTTCCTGATT CAGAAGGTGGGACTCACAATTGTAATTCTGCTAATGGTTGTCTTTCAGTCTATCAATTGC TTCATAAATGCATCCACTGTTCCTTCTTCTTCTGCCCTGCTTATAATTTTCCATGAGTCC ATATATCTTTTACACTGTCTTTAGTCTTATTCACTAAATTAAAAACTAATTTTTGATAT TTGGTATTCATGACAAGACAATTAGTAGAATTTTGATGCTTCTTGTCTGCAATTACAGAA TCAATATATTTCTATATTATTGTATATTCTCTAAATCTTATTTTGTATAATAGCTTTCA GCATGTTCTTTAATTCTGTTTAGATATTTAGAAAGTATTTGTTGTTATTCTGTAATTTAT TTCAATATTCAATTATAGTTTAATATTTTGTTATCTAGTGTTGTCTTGATTTTGATATAC GTACTGATTTTGTAGATCCAAATTCCTCTTTCCTATCAGAGAATGCAATTTTTTACTTGG ATAAATAAGAATCATATCTCCTCTGCTTGCTACCGTATTGCATACATTCATGGGTAGAGA AAGAGTTAAGCTGATGAGAGTAGGAATTAAGGTAGACCTGTTTGGTAGGTTCTCCCAGAT TTCAGAGGACAGACATCTTTTTTTCCCTGCCTTGGTCATTTAAACTTTTTGGATTTTGGA TTAAGTGTAGGCAGGGAAAATGTATCAGATATTTTTATTTTTCTTTGGTGCCATTTGTCC TTCTCTGCTTTAGGCAGAGAAGCATATGTAGTCCAAGAATGTGCTTTTCTATCCAGCTAC ATCAATAATAACAATTAGTAAAATTCTACTTAAACTTAGACCTTTGCTGTTCTCTTTTCT CTGCTTGTGTTAAGTCATGCTCATGATTCTGGCAGTTTTCCACAGTACCATGTACAGAAA GCTTGAATAAGGTACATCTAGAATACTCATATATGTTCACTTCAAAAACACATTTTTGTG GAATTCTAAATGCAAATCTCAATAGTGCAATTCTAATTTACAATGAGAAAAAACTAAGGG ATTTTTTCTGGTGATTCTTTTTGCTCATTTATAAATATGTTTTTAAATGGTAAGCAAATA TATAAATTAAGCTTTTCCTTACGTAGCTACATTGATTTACTAGTGGTGGAAAAGGTTAAG CAAAACTAATTTTCATGAGTGTAAATGAATTAGTAAGTGACATATGCAATGCTTAAGGGG AATTTGCATAAATCTATGACTGATACTCAACCTCTTGCTTAGCGAGAAGATAATTAAAAT ATTTTATACTTCAAGAAGACCTAGTTTTCCAAATTATTTACATCCACAAACTCAGATTTT ATAGCAAGTAAGAAAAGTTAAGTCAGAAGCATATACTATTAACAGCTACTTACATTGCTC AAATTTAATATACGATTGCTGCTTTTGTTGGTTTTGAAATGTTTCTTGACCATGGATCTG GATACAATAATATTAAAAAATTAAATGTAAGTTCCTGCACTCACAGTAGAGGTAAGTTCA AGGTTATAAGAGAGCTTATAGATTCTGAGATTTGGAAAGAAGAGAATAGAAAAACTTTT CAGATTAAATAATGTGTTAATTGTGCTTCTAAAACAGCTTTGGTGATCTTAATAAAATAA ATATTGTTTTTATTTCCATTTTTGCTTTTCAGACAAGAAATGCTACTTGATGGCTGCATA TATTTGTTTTGTCTCTTTTCACCACCTACTCTTGCTAAATACTCTCAACCCACTCATGAA ATTAAAGCACATTGGAAAACATTTATCAACTACCTGTAAATACAACCTATGCTCTTTT GTGGAGGTGATAGACATTCATCAATGGAATAGTTGATCTAAATCCTAGTCTTCATTATCT TGTTTTATACATTCTTGTCTTAATCAGTTTGGGCTGCTCTAACACAATACCATAGACTAG GTGGCTGATGAACAACAGAAATTTGTTTCCGACTGTTTTGGAGACTGGGAAGTCCAAGAT CGAATTTTATGTCTGGTGAGGGCCTGTTTCCTAATTAATAAACATCTGTTGTCTCATATG TCCTCACATGATAGAAGGGGCAAAGGAGCTCTCTGATGTCTCTTTTTTAGAATATTAATC TCGTTCATGAAGGCTCTGCTCTCATGACCTATTCCTTCCCAAAGGGCCCCACTTCCAAAGA CCATCATATTAGGGATTAGGTTTCAACAAATGAAGCCAGGGGGAGGTTGGTAAACATTCA ATCTATAGCAATGCCTATCTCCAGGAGCTGCCTGTGGAAACACTTTTATCTGATATGGTA GTTTAAAGCATGGCAGGGATAAGTGGTATGAGGAAAACTCTCCCTGCCACCCAACGCACA CATCCCACTTAAGCTTCAGCAGCTCCAATTTTATCTGTGTAATATTTGGTTCCACATCAA AGTTGTTTTGAATATACTTCCATTACCTTAAAAAATGTAAAAACACTGCTTTAAAAAGCC AAGCCTATTCCCTTTTCATTATTCAGAGTTCTTCCAGTTTTACCGTTACATCAAATTAGA ACTACATAATTAGGAACCCCTCTCTAAATTTGCCTCTATACAGAGAAAAACTGTGCCTGA GGACAGAATTTGTACTTCGTTCCATACATAAAAACTCATTTGACAAATAACAAGCATAGC TCCAAGCTCAAAGAATAGCTTAATTTTTCCTGATTAGTTTATATCTCTCTTATTAATCAA TGACATTTAATATTACAACCATAGCTTGGGGTTTTAGTTTATTTGCTTTCTATCTTTTTT ATACTGTCGGCCTACCTGTGCCCAACTATGTTATAGTCAGGGGTTGGTAAAATAAAGACA AAACAAATCCTGTCTTCCTGGAGATCACCTTCACTGGGGGTTGAGAAACAATAAGAACAA GTAGTAAGTAAAATATGTACATTAAAATTTTAGATGAAGTTAAGTGCTATGGAAAAAAGT AAAATGGAAGAGGTGTTATGGAGTACCTGTTCGGGTATGGGTTCAATTTACAAGTGGATG GTCACCTTCTCACTGATAAGGTGACATTTGAGCAAAAGTCTTCAGCAGGAAGGGAGAATG CCATGCAGTTATCCTAGGAAAGAACATTTCCAATATAAGTAACAGCCAGTGCAAAAGCCC

FIG. 1P

TGATGTAGATGCATACCTTAGGTATACGAGTAACAGTAAGAAATTAGTGGCACGAAAGAC AGATGTACTTGGAAACCAAAAAGAATCTCTGGTAAGAAATTGTAAGTCATTGTAAGGACT TAAGGTTTTTTTTTTCCTCCCAAATGAGATGGAGATCCATTAGAAGGGTTTGCGTAGA GAAATAATATGATCTGACTTATATTTAACAGGACTACTCTTTTGCTGAATTGAAAATTGT CTCTAAGGGTGTATATCAGATCTTATATTGATCTTACCCTTCTCTGTTCAATATTTAACA ATCCCCTCTCATAAGGGCTGTGAGGGCCTAATCTGCTTACCTATCCAGCAGGCTGGGAAT GACACAGAGCACTCACCAGGAGCACTCTCAACCTATGACTCATGGAAGTTGGTAGATGAA TACCCCAGCTCTCATATTCCTTGGGTGGAAGAGCTCTGAGATGTGTGTTCTACACCATTA CCCAGAGGGCACCCTCTGGATTAGGCTCAAGTTGCTGACAGTAGTATCTTGCTGACTAAC ATAATTTTTATTAATTTTCTCCCCATTTGACCTTATTTCTCCATTTTTCTAATAGTGTTC ATTGGTATCACTTCCAAAATAAATTACCTTTACTTGAATATTTTTCTTAGAATCTTCTAT ACAAACCTGAGCTAATACTGGGGCAAAGAGTGGAAGCAGGGAAATATTTTGTAGGTGTTG TGGTGATGTAGGACAGAGCCTGATAGCTTGGATCAAGGTGGTAGCAAAGGAGATTGTAGA AGCTATCACACTCTTTATATATTTTGAAGACACAGCCAAGAGGTTTGGTGGAAAAATGGA TTGTGAGAAGTAATAAAAAGAGTGGGAGAGAAAGTCAAGGATGTCACCAAAGTTGTCCTA AGCAAGTGGAAACTTAGATTTGGGAGAATCAAAAATCCTAAAATATCCAAATCCTCTCCC $\tt CTGCCTTCCCCTCCCCTCCCCTTCCCTTTGGAGATAGGGTCTTGCTCTGTTTCAC$ AGGCTGTAGTCTAGTTTCGCGATCTCGACTCACTGCAGCTTCGACCCCCTGGGCTGAAGT AATTCTTCTACTTTAGCCTCCCAGGCAGCTGGGACTACAGGATTGCACTAATGTGCCCAG CTGATTTTTTTAGTTTTTTTTTTTTTTTAGTGGAGATGAGGTCTCGCTATGTTGCCTGAG CTCAAGCAATCCACCCTCCTCAGACTCCCAAAGTTCTGGGATTACAGGTGTGAAACACTG TGCCTGGCCCAACATTTTATTTTCAAATATTTAAGTTTTGAATGTCTATTCGATAACCAA GTAAAGAAGTCAACTAGAATATATGAGAATGGAGTTTTCTAGAGAAGTCTGGGTTGAGGA TGTACTTTTGGGAAATGGAGCACATACTTGGTATCTAAAGCTGTGAGCCGAGATGAGATC ACTAGGTAGGTAAATATAGATAAATTAGAGAAAATATCTAATAATTGAGACATGGAGTAC TATCATAAATTTTGAAAAGACAAGAAAATGTGAGAGATCGAGAAGAATGGCTGGGGAAGA AAATGCTACTGATAAGTAAAGTGAAATGTAGAATGAAAGTCAACCATAAAATTTGGCATT ATGGGGATCATTAATGACCTTAAAGAAAGTGCTTTTAGTGTAGTAATAGAAAGATGCAGA AAGTAAGTAGAGTGAATTCAAATTCAACAGAGAATAGACAGAGAGGAATTGAAGACATTT GGCAAGTGTTTGGAGGCCAATTTATACTCAAGAATAATTTCTTGAGTTGGTTTTTTGTGT $\tt TTGTTTGTTTTGATTGGTTAGTGTGTTTATTTTTTAGACGGGATTGGAGAAATACTTTC$ ATTTGTGTTTTTACCCATGTTTTCAGCCTTGCCCTGGCTGCCTGGTATAACGCAACTCTA $\tt TTTGTTATTCTGCTATTATAGTTTCCCTAGCTTGAATTTTTTTACACCCTTATTATAATT$ GTAGCGTTGCATGCCTATTTCAAAACATCTCATGTACCCCATAAATATATACATCTACTA TGTACCCACAAAATTAGAAATAAAAAATTTAAAAATTATGATTTTTTAAAATTTGTTA TAGTTGAAGGAAATTTCAGATATTTTGGTAGCAGAAGGAACTGAGTTATGGCTCAAGAGT TTTTTAATAAGTGTGAGTGGAGTTATACAAACTACTCATTAAAATCTTTATTTGAATTTG TAATATCTGAAACCATTTTCATATTGAAGAATCACTTAAAATAGTCATAAAATGTAAAAT TGCAAGACAATTAAAAACAAAAATATGATTTCACGACTGTGATAGTACCTGAGAAATTTC TTCATCTCCTTAGTAAGAGAAGTATTACACCTATTTATAGTTATTTTATGAAACTAGCTA AGATGAATTATGTAGAAAAGATACAGATTTTCAAACAGAAACTAGAATTAATGGAAGCTA AGCGAAAAATAGCATCTACCTATAAGGATTTGCAAAGCCAGTAATCTTTCTAAAAATATC AGCAAACCCAGAATTAAGGCTTATGTTCTTAGCTCATTGTAACTAGATCAAAAATAAAGA AGGCCAAATAAAGGTATGTGACATTTGTTGAAAACCTGAAGTGTCCTATATGCAGAAATA TTTTTATCATTAATTAATTTCAGAAACTTCTTAACATGACATGATCCTCTTGAAAAGAT CACATCAAAAAAGGCAAAATAATTGCATAATTATTGTAGAATAATTTTTGTGTGAGTATT TTTAAAATATGCCTAATTTTCCAGGCATTGGTTTGCTTTGCTATAAAATGGGAGGATAGA AAATAACTTTCAAAATATCTTATAAATCTAAGAATCTTTGCATCTTATAAATCTAAGAAT CTTTGGAAATTCATAGATTATTGAGATGGAGTCTCGTTGCTATGCATTGTAGCAAAGTTG

FIG. 1Q

18/51

GAAATAAATTCTAAATTTTATTTCATTTATATTGATCAATAAATTGTTACATTTCACTAA TACAATAAGGAAAATTTATTTTACCTGAGTGTATGTCTAGCTTGTGAAATAAAAATGCTC AATTATGAAAGCATTTATTGCCATTTTGAATGAAAAATGTAATATGTAGAACAGAATTTT TTTTGCCTTGAACTCAGTTAAATGTAGAAATTGATAAGGACTTGCATTTTCATGAACTTA ATAATTATCTGTCTTTTCAATGGTCTCCATATCAAGTCTGAGAAATATGGATGTGATTTA TTTTAAACCTCACCATTTGAAGTAAATCTAAAGATTCCATTAGGTTATGAGCATATAGGA TACAAGGACCATATTGACAGTTTTGTGGGATTGTATTAGGATAAAAGGGTAGGAACAATG GGGAGAAATTATAGCTTACAATAGGGAAGAACCAAAAATTGTTGCAAAATGATGGAACA GGCTGAAAGAATGATATAACCTCCTAAACACTTCAAATGTTTAAGCAGTTCATTGTACCA GTTTCTTCTAACGATAAAATAGCACAACTCACTTTTTTTCTAACCTCTAAGAGTATATTA TTAAAAGTCTGCTACAACTACAGATAGAGGAACAGTTTGTAGTATCCGTGATCCTAGAAC **AAATTTAGCTTTTAATATCTTGTCAACTTTTTTGTTTTAGTATCTCTTCCTTGGAACTAG** CTGAGCTTTAATGGCATCATCATGTGATATGACTTGAGATTTATATTTGGAAGAGCTTTG AAAAATCACGGATTGTTACCCTAATGAGGTGTTATTCAGTCTTTTAAACAAGAGCAATTT ACTTGGCTGGGCATGGTGGCTCACACCTGTAATCCCAGCACTTCGGGAGGCAGAGGCTGG TGGATCACTTGAGGTCAGGAGTTTCAGACCAGCCTGGCCCAACACGGTGAAAAACAGTCT GTAATGCCAGCTACTCGGGAGGCTGAGGTGAGAATCACTTGAACCTGGGAGGTGGAGG TTGCAGTGAGCCAAGATTACACCATTGCACTCCAGTCTGGGTGACAGAGCGAGACTCCAC CTCAAAAATAAAAATAAAAAAAAAAAAAAAGAATGAATTGCTCATAAATGTGCCTCACTGAT GATTAAATTTAATCCTGCAAGATTATGTCTTTTGATGGAAATGAGAGGGTTTATACAAAG TTTTATTCGTGATGTTATCTATGTCATCTATTGATTTCTGCTCTGATTCATGTGGATGAA GTTACACCTCACACTTTAAGCTGGTGTCAGTCTTCCCATTTTCTGCTGTGATGTGTACTC AAGATCTCCAGATTACATCTGTAATGTAATGCAGCCATGATTGTTTATAGGTACATTTAG ATGAATTCAATGATGAGTTATGTTGTAATAAGTGTCAGATTTAGATGAACCATACAAATA AAAGAACCATGCATTAAAATGACAAATGTGTAAAAGCATTATTTGGGCCTTAAGTCAAGG CCCAAATGTGGATACTGGTACTGAGACATCTTTCAGAAAGGAGGTATGAAGTACTGAAAA ATATTTACAAAATGAAGACTACTTTTATCTTACTTATCATGATTCTTTATTACATATGC ATTTTCTAAGATAACTATAGTGCATTAGTTTGTACTATGTTAATAATAATAATAGGGTAAA TCAAACAATGTTTTCTAAATCCATTAAAATAGAGTTCCCTAAGGGAGTTAAAACAATTAC GTTCTACTGTATATTATTGGCATGCTTCAGGAGACATGATTTAATCTCTAGACTATCAGA ATTCAAGAACTAGTGAGTCATATAACAAAGGAGGCTTAATCATGCCATTTAAGTGTCATG GAAAAAGGTTTATTGGTCAGGAAAAATTAATTAGAAAAAAGTTATAAAATACTTCACTAA GAAAATAAAATGTCAGGAAGCCCACTTAGACAATGAGTGAAAATGAAACAAATTCAAGTT TTTACAATATTTGGTTTCTATAGGATTGCTTCATTGTTTTTGGTTTTTTCCCCATA TTGTTCACATGCCACAGACAATCAATTATGAAGAAAGGAGAGACTCGTAGGAGGCAGG GCCAGGCTGTTCACACTTTTAAACTAGGTAGCCACAAATGAGGCTTAGTTACAAAAACTT GAAAACTGGATTCTTCCCAATGTATTATACATCCCCAAAGAAATGATGAAGTTCCTTACT TAGGTTTTCATATTGGCTTAGATTTTTTTTTTCATTAACTTGCAATTTGTGGTTGGGAAAT GATCTGCTTTTTGTTTCAGGTTGTTTAATGTTTTCCAATGTAATATTCTTCTTGCACTCC AGTGAGTTTATTTACAAAACATTTAATGTCATTTGCGTCTTCGAAGAACAATGTATTCGG TTAGAACAAAAGTGAGCTCCTGCATAGAGCTTATGATGGTTTATAATTGGTAAATTATTA CTCATTCAAAGGGACCGTTCACCCACAAAATGCCTTTTTGTTTATCTTTTGGAATGACAC CATTGGAAACTCAGTATGGCCACTTTTATGGTAATAATAAAAGTCATATATAAAAAGGAT CTTTAAACTACTAAATACAAATAAATTAGTAGTACAGTCATTAGGATTGCTCTTAGTTTG TTAGTGTTGGAATAGACTTTTGGATTTTCTTCCTAGCTTAGATTGATACAATGTGATGGG GACTTGCTCCAAACACAGGAATAGGTGGCCTGCAGACACACTCTGTGATGCTGTAATT CTAATCCTCACTGAATATATCAGGGGTGGACATCTGGCCTGGGGCAATTCAGATACTTTT TCTTAAAATTTATACTACAAATTCAAAAGTGGTAACTCATCTCTGCCATCACTTATAGTA

FIG. 1R

TGGAGAATAAAATAAGTAAATTAGAACAGGAAAAATGCCAAAACACACAGACATGACCCT GATAGTTTTCCATTTCCTGATCACTGTCCCTTCCTGTGGCTGGATAAGGAACTGTCTCTA GGCTCTGTAAGACATATTTGCATCCTTACGACAAATTTCTACTCCTTTTCATAAACTAGA CTTGGGTTCTTTAACTTGCAACAGCAACAACAATAAACGATTTTGTTGGGTACAATCTGA TTTTATTAACTTCTGGATTTAAAAGCCCTTCTAAATGTTGATTGGCATTGTTTTTACTTC CTAAGAGTACGCTCATGCACCACATAGTGATGTTTTGGTCAACGACAGACTGCATTTACG ACTGTGGTCCCATAAGATTATAATACCATGCTTTTCTGTACTTTTCTATGTTTAGATATG TTCAGATACACAAATGCTTATCATTGTGTTATAATTGCCTACAGTGTTCAGTACAGTTAC ATGCTGTACAGGTTTATAGCCTAGGAGCAATTGGCTATACCCTATAGCCTAGGTGTGTAG TAGGCTATACCATTAGATTTGTGTAAGCATACCCTATGATGTTTGCACAATGATGAAATC AGAAGTACAATAACTTTCAAATCCTGAATGTTCTGTACTTTCCATCTCACAAGCATTTTG CAAAGCATCAAATGGTATAAGCCAGATTACTGTTAAGGCAACTTGGAATTAATATGCTGC TCAGTTCTGGAAAAGGCATATTCTGTAAATATAGATGAGAGAATATAGACTTTTTCCCTC TCTTCTTACAATCCACATTCTATTCAGTATTTCATTTACTTGAGGGGTTATATGCTACTT ATCTTTATCTGTTGTGGAGTGAGGACACATTCCAAATGCCTTGGTATTATTAAAAGCCCT TCATGATGTGGCCCCATCTTTATGACTTTTCCTTTTCAACTGTGCCCTCTAGCCTTATT TGATTTCTCTCAAATTCTTAAACACAGCATGCTTCACTGACCTTTAAGCCTTTGCACATA CAGTGTTGATGTGGAGCTTCCTGACCAACTCCTAATTCTCCTTCAGGCCTCAATTTAAAC ATCACTTCCTCTGGGAAGCTTTCTATTATTCCCAAGGTACTGGGATATGTTCTTGCACAG CATGCTGGGCTAATGTCACAATGGCTACCTTGTTTTATTGTTAGTATTTGATCAGCGACA CCTTGCCAGGGAGCCCCTGAGTATTGTCTGAGCAGAAACTATGGCTATCTTGTCCCCTGT ATATCTCAAGATCATGCTGAAAAGCCAGCATTCATGAACAAATTCCTGTGCGAAGATTGA GAATGAAAGATGAATAAGAGGTATCTTTAGAACCCAATTATGGCTGCCGTTGTTCCCTGA GTGTGAGGCTTGCTGTTAGAGTGACAGAAGGAATTTTGACTACTCAAGACCATACAAATT TGGAAATGACTCCAAAGTAAACATGGTTAGATAACTACACATTCCATTCCCCCTTTTTTA TTTCTATAGAATCCCAACTTTGTTCAAGTAGTAACATGCCCAGCTTCAGAAATGAGTCAT GATTTTTCTAAAGCAACAATATCAATCTTCTTTCCCTTCCCCAGTGATTGGTATGGAAGT TTTTGCTTTCTGCTGTAAATCAAAAGCAGAAACAGGAGAAGATTCTTTTGGGCCTCTTTC CCTCTTCCTGGCGTGGAAGTAGTTGTGAGAGCATATGATACCCAAAGTTTCGGTAGACAT TTTATAATTATGTGATGAATAACCTAAGGATAATTAAACATATAAAAGAATGGAGAAAGA GATTAAGTGATGTTATCTTCCTTTAAGGCAATCAAAATGCATCTGACAAATGGCCATCTA ATTTAAAATTCCAACTATGTAGACATCTCAAACAAGTCAGTATCTCAAAAAATATACTA CAAAAATTCTCATGTGTCCATTGGGGATAACTTCCAATGCTCTTTCATTGGTATTGTAGC TATGGCATTTGATTTCCAATTGTATGTGGATCAGGTAGTTGCAGGGTGACTCTCAAGGGC AACAAGACAAAGTCAAACCCTAGGTAGAAATAAGAAGGAGCTAGTACAGAAAGCAAATGC CTAAGGTGTTGGAGAACATAGAAAGGTAGAGTGGAATGAAAAAAGAAAAAAACACTAAATA GCAGCACATAGAATCTTGGGGTTTCAGGGATATTGTTTATGAAAGGTTAGAATAGGCAAC AATTACCTGTCATATGGGTCTTGTCATTTATTTATAATTTAAGGAGAATTAAAACTGAAC TAGTTGCTGGGGAGTGACATCAGCAAGATGGAGATATAGAAATCTTCAGGACCTCCTTCC GTCCATGGAACCACTGACTCAAAAATGACAAATGGAAAAAATTTACTTTCTGAGAAATCA AGAAGCCAGTTAAGAGGCTCCTGTATCTCAGATGAGTGCAAAGCCAGCTGCAACAGAGCC AGCAGAAAATTTGTTGTACTCACTCTTCATGGTCACTTCTGGCATAGCACAGTGCAATCT AGAAGAAATTCTCGGCTCCTGACTACTTTCTTGGAAAAGAAAAGAAAAATGTACCATAT GTCTAATATTCTGATGGGGATGGGGTGTGGGCTGCTCAAAGGACTAGCTTCCGTCATGCC TAAATACAAGTGCTAATTGGGAAGTCCACAATGTTGGGGGCTGCAGAAAACAAGGGCAAC AGTTTGGACTAGCATGCACTCATTTGCCGCAGTTCCTCCTCTCACTTCATAGAATGAGTA GAAGAACCCTTAACTCTCAAGGTTTTTTTCCTGGGGAGAGAAAGAGTCAAAGCAATTATA

FIG. 1S

20/51

ACCAATCTCAGAGTGCAGATGGAACCTAGCATATTCTAGATGCCTGGGGGCCATTGAGAA CAAAAGAGAGCTAGGCAACTTTCAGCAGCTCCAGAAGAACTGTGGTACCACAGATAGACA AGAATTTACACACACTGGTACAGATAAGATGAATTTGCAAAAAAAGAATAGAGGCCCCAG AATTTCTAGCTGGGTTTTTTGGTGAAGGCCTTTCTCTGTATCAAGCTAGTCCCTAAAGAC TGGGTGAGGTGGTTTTTGTTTTGTTTTACATTTTTATTTTAAAAGATGGGGATCTCACTTT GTCACCCAGACTTGAGTGCAGTGATGCAATCATAACTCACTGCAGCCTCAAACTCCAAGG GTCAAGTGATCTTTCCACCTCAGCCTCCTGAGTAGCTGAGACTAGAGACACATGCCACTG TGCTTGATTAATTTTTTTTTTTTTTTTTTTTTCGTAGAGATGTGGTCTCACTTTGTTGT TCAGGCTGGACTTGAACTATTGACTTCAAGGGATCCTCCTGACTCAGCCTCCCAAATCAT TGGGATTACAGGCATGAGCCACCATGCCTGACCTGTTTTGTTTTTAAAAAACTCAG AAAAATTTCAAAATAGCAATTATAAAGACAATGAGCTTAGAAAACCAATTAATGGACAAA ATATGATAACCAAATTGAATATTACATTAGAGGAGTTTAATACTAGATTTGAACAAGCAG **AAGAAGACTAAAAAAGAGTGAAGAAACCCTAAGGACATCATCAAGTAGACCAATATGTGT** TATCAGAGTTTTAGAAGAAAAAGACAGAAAAATAGGCATAAAGCATCATTGACAAAATAA TGACCCAAAACCTCCCAATTATGAAAGACAATAGATATTCTGAATCCAGAGCACAATGGC CTGCAACTAAGATGAACCCAGAAAAGTCTATACTTCAGCACATTATAATCTAATTATCAA CAAGGGCTGTCATGAGAATATCAGCAGATTTCTCAGCAGAAAACTTGCAAAACAGAAATA AGTGGGATTACATATTCAAAGAGCTGAAAAAAAGTCTGCCAACAAAAAATCCTTTATCCA GAAGAATTTTCTTCAAAATGAAGGAGAATAAAGGATATTCCAGATAAACAAAAGCCAAGG GAATCCATCACAATTAAACCTGCCTTACAAGAAATGCTAAATGAAGTTGTTCAAGTTGAA ATAAAAGAACGCTGAACAGCAACACAAAAGCATATAAAAGTATAAAGCTCATTGGTCAAA GATAGATATAAAGGAAAAACAACGGGATATTATAATGGTGGTGGGTAACTTACTCTTCAT CCTGGTATAGAAGTTAAAAAAACCACAAGTATTAAAATAACTGTAACTATAAAATTATT AATGAATACACAATGTAAAAATATGTAATTTGTGATACCTGATAACATACCATGTGTGGAG GGGAGAAGTCAAAGTGTAGAGTTTTAAATAAGACTGAGGTTAGGTTTTTATCACCTTAAA ATAGATTGTTATAATATGTTTGATTTAAGCCCCATGGCAACTACAAAGAAAATACCTACA GGTAATAAACAAAAGAAAATGAGAAAGAAATGAAAGTGTGTCTCAGTCCATTTTTATTTT TTCTGGAGGCTGTGAAGTTCAAGACTGAGTTGCTGCCTCTGTTGAGGGGCCCTTCTTATTG CATCATAACATGGCAGAAGGCATCACATGACAAAAAAGCAACAGCAAGAGCCAAACTGGC TTTTATCATAGGCCTAGTTTGTGACACCTTACATAGTCCTATGAAAACCCATTAAGCCAT TAGCCCATTAATCCATTAATTCATGAATAGATTAATACATCCATGTGGGGAAAGCCCTCA TGACTCAAACCTTTCTCAAAAAACCCATCTCTTAATACTGTTACATTAGTATTAAGTTTT AACATGAGTTTCAGAGTCTAGAAATATTCACACCATAGCCTTTCACCCATGACCTCCCAT AATTTATGTCCTTATCATATGCAAATACCTTCATTCCATTCCCGTAGCCCCGAAGTCTTA ACCTGTTCTAGCACCAACTCTAAAATACGAAGTCAAGAGTCTCATCTGAGACTCAAGGCA TGATCCATCCTTGGGCAGGTTCCCTTTCAGTTGTGAAATCAAAACAAGTCATATAATTCT AAAATACAGTGCTGGTACAGGAATAAGACAGACATTCCCTTGTCGAAAGGGAAAATAAAC TAGAAGAAGGGGTTAATGGTCCCCAAGCAAGTCTTTAACACAGCAGGGCACATATTAAAT TGTAAAGCTAAAGAATACTCTTTTTTGGGTCCATGTTAAGCATTCTCTGCACAATGTGGG GAACACATTGAGCCACTCTGCCCCTATGGCTTTGCTGTGCTCAGAACACACTTCAGCTTT CTCAGATTGGAATTGCTCATTGGTGCCTGCAGCTTTCCCAGGTGGGCACTGCACACTGCT GGTGTTTCTATAATTCTAGGATCTCAAAGGCAGCTCTGGCTCTCACCCCGTATTTTTACT CAACATTGCTGTAGTGGGGCTCTCAGCCATGGCTCTGTCCCTGTGACAAGTCTCTGCCTG GGTCCCCATGCTTTTAGATACATCCTCTGAAGTCTAGGTGAAGGCCATAGTGGCCCTACA ACTCTTGCATTCTGTATCCCTGCAGAATTAGCACCAGGTGGACACTGCCAAGGCTTATGG CTTTTGCTTTCTGGAGCAGTGAGGTAAGCTACACTTGGAGCCTCTTGAGCCAGTTGGAGT GGCTGAGGAATGATGCGCTCACATGAAGGGAGCAGAGGAGTCCTGAGCAGCCCTGGGCAG CAAGCTGTGGAGAGTACCCTGGGCCTGTCCCCTGAAACTATTCTACCCTCCTTGGCCCCT GGGCTTTTCATGAGAGGGGGCAGTCTTAAAAATATGCAAAATACTTTTCAAACATTCTCC TCATTGTCTTAATGAATAACATCTGACTCCCTTCTATCAGTGCTAATCTCTTTAGCAAGC AGTTTTGCTGTTTACATGGCTAAGCAAGCTGCAAACTTTTCAAATCATTTTGCTGTGATT CCCTTTAATTATACATCTGTCTTTAAGTCATGTTTTTGCTCCTGAATTGGCCAAAAGTAA

FIG. 1T

21/51

CCACACAGCCAAAAGTAGCCAAACAGCATCATGAATGCTTTGCTCCTTAAAAATTTCTTC TATAAGATATTTTACTTTATTATTGTCAAGTCTGGCCTTCTACACAGCCCTAGAGTATGG ACACAGTTCCAGTAAGCTTTTTGCTACTTTATACCAAGTATGACCTTTATTCCAGGTTCT GATACCTTGTTCCCCCTTTCTGTCTGAAACCTCATAACGGCCTTCATTGTCTATATGTTT ACTAGTATTTTGGCCATAATCACTTAAATAATTTATAAAATGATTCAGACTTTCCCTAGT CTTCTCATCCTCTGATCCTTCACCAGAAGCACCCTTAACACTCTATTTACAGCAATATAA GATTTTTTTTGCCTGCTCCTCCAAACCCTTCCAGCCTTTGTCCATTACCCATTTCCAAAG CCACTTGCACATTTTTAGGTTGAGCATCAGCCTCACTTCTTGTTACCAAAGCCTGTATTA GGGTTCTCCAGAGAGACAAAACCAATGGGATATACAGAAGGGGATTTGTTAGGGAAATTG GCTCACACAGTTATGGAGACTGAAAAGACCAAGGTCAAGGGGACGTATCTGGTGAGAACC TTCTCATTGTATCATAACATGGCAGATGGCATCACATGCTAAAAGAGCAAGAACAATAGC CAAACTGGATTTTATAACAGACCCACTCTTGACGACTATCCTATTCCTGTGATAAGCCAT TAATCTGTGAATCCATGAGTAAATTAATCTATTCATGAGGGCTCTGCCTCTATTGTCCCT TAAAGGCCCCACTTCTTAATACTGTTACATTGGGGATGAAGTTTCAATATGGGTTTCAGA GGAGACAAACATTCAAACCATAGTGATGTCACTACAAAAAATTAATGAAACACAAAGGA GTACAGTAAGAGGCAAAATACAGATAAAAGTGCTATATGATATATAGAAAACAATAAAA TGGCAATAGTAGGAGTTTATCTGTCAGTAGTTACTTTAGCCATÁAATGAACTAAACTCAA ACAAAAGACAAAGATTAGCTGACTGGATTTAAAAAAATACTATATGCTGTCTACAAGAAGT ACAAGGAGCCCACTCCAAATTTGTAGACACACATAGGATAAAATTAAAAGGATGGAAGAA AGTATTCCATGTGAATGGTAACCAGATGAGAGCAGGGCTCATTATACTTATATCGGACAA ATAAATTGTAAGTCAATAATTGTCACAAGGAACAAAGAAGGACAATATGTAATATTAAAA GAGTCAATTCACCAGAAAGATATAACAATTTTAAACATATATGTATTCAATCTTAGGGCT TTAAAATATATAAACAAATATTAATGGAACTGAAGGGAGAAAGACAGCAATACAACAATA GTAGGAGATTTTAATTCTCAGCTTTCTTTTTCTAGAGACAGAGTCTCACTCTGTCACTCA GGCTGGAGGGCAATGGTACAATCTCAGCTCACTGCAATCTCCACTTCCCAGACTCAAGTG ATTCTCCCACTTCAGCCTGCTGAGTAGCTGGGACTGCAGACATGCAACACCATACCCAGC TAATTTTTTAACTTTTTGTACAGATGAAGTCTCGTATATTGCCCAGCTGGTCTTAAACTC TTGGGCTCAAGTGATCCTTCACCTGGGCCTCCCAAAGTGCTGGGATTATAGGCATGAGCC ATGCTGTGTCACAAAACATGTTTTAACAAATTTAAAAAATACAGAAATCATATCAAATATC TTTTCTGAACACAGTGGAATGAAACTATAAATCAATTATAAAAGGAAACTGGCAATTTCA CCAATATGTGTACATTAAACAATAAATTCTTGAACAGTCCATGAGTCAAAGAAGAAATTA TAAGGGATATTTGAAATGTTTCAAGATAAATGAAAATGTCTCAAGATGAAATAAAAAGAC AACATATCCAAATTTATGGAATGCAACAAAAGTGGCAAGAGTTAAGTTTATAGTGGTAAG TGACTACATTATAAAAGAAAAAGATTTTAAGTAAACAACCTAACTTTACACCTCAGAAG CTTTTTAAAGATCAATAAAATTTACAAACCTTTGGCTAGAATAACTAAGAAAAAAGAGAG AAGACTCATAAATAATATTGTAAATAAAAAAGGAGCTATTGCAATCAAAGAGGCAGGAAC AATAAAGATTTTCAGGCTATTCTGTATAATTATACACTAACAAATTGGATAACCTAGAAG AAATGTATAAATTCTCAGAAATACACAACCTACCAAGACTGAATCAAGAAGAAATACAGA ATCTGAACAGATCTGTAACTAGGAGGAGATTAAATCAATGATCAGAAACTTCCCAAAAA AGAAAATCCCAGGATCAGAAAACTTCACTGGAGAATTCTGCCAACATTTAATAGAAAAAA AAATGCCAATTCTTCTCAAACTTTTGCAAAAATTGAAGAGGACGAAGCATTTCAAACTC ATTTTATGAGTCCAGCATTTTCCTGATACCAAAATGAGATAAAGATATTACAACGAACAC ACACACTTTCAAACAAGCTACAGGCCACTATCTCTGATGAATGTAAATGCAAAAGTTGTC AATAAAAATAGCAAACTGAATTCAACAGTGCATTAAAAGGATCACACACTGTGACCAAG CTATATTAACAGAACAAGGGATAAGATCACATGATAATCTCTATAAATGCTGAACAATCA TTTGACAAAGTTTAATACCCTTTCGTAATAAAAATACTCAACAAACTATGAATAGAAGGC ATGTACCTCAACACAATAATAAAGGTCACATATCAAAAGCTAACAGATAACATCATACTC AATGGTAAAAACTGAAAGCTTTTCCTCCAAGATCAGGAACTAGGTAAGAATGTCCATTCT TGCCATTTCTCATCAACGTATTACTAGAAGTCTTTGCTAGAACAATTATGCAAGAATAAG AAATAAAAAGCACTGAAATCAGCAAGGAAGAGGGAAAATTATCTCTATTCCCAGATATAA TAATCTTATATGTAGAAAATTCTAAAAATCACACAAGGAAACTGTTGCAACTAGTAAGTT

FIG. 1U

CATCAAAATTGCAGAACATAAAATCGAAATGCAAAAATCAGTTATGTTTCTATACAATAG CAGCAAACTCTCTGAAAAAGACATTACAATCCCACTTACAATATTATCAAAAATGACTAA AATGTTTAGTAATAAGCTTAACCAAGGAGGCTAACGACTTATACACTGAAAACCATAAAA GCATTACCAAAAATAATTTTAAAAGACACAAATAAATAGAAAGATAATTCTGTTTTCAT GGGTTAGAAAACTCGATATTGTTAAAATGTGCACACTGCTGAAAGCAATTTATAGATCCT ATACAATCTTACCAAAATTATGATGTCATTTTTTTCAGAAATAGAAAAAAATCTGAGAA CCATGGATACTTAGAAAATCTGGAGAAAGAAGAAGAAGTAGAGGGTCTCATGCTTCCTG ACTTCAAAACATATTCCAAAGCCATTGTAATAGAAACAGTTTAGCACTGGCATAAAGACA AAATAATATACAAAGCACAAAGACTATGGACAGGATAGTCTCTTCAACAATTGTGTTGGG AAAACTAGATAGCCATATTCAAAGGACTGAAATTAGACCCTACTCAAAAAATCAAGTCAA AATGAATTAAAAATTAAAGATCTGGGCCGGGCGTGGTGGCTCACGCCTGTAATCCCAGCA CTTTGGGAGGCCAAGGGGTCAGATCACGAGGTCAGGAGATCGAGACCATCCTGGCTAAC ACAGTGAAACCCCGTCTCTACTAAAAATACAAAAAATTAGCCGGGCGTGGTGGTGGGGCGC CTGTAGTCCCAACTACTCAGGAGGCTGAGGCAGGAGAATGGCGTGAACCTCAGAGGCAGA GCTTGCAGTGAGGTGAGATCACGCCACTGCACTCCAGCCTGGGGGACAGAGCAAGACTCC ATCTCAAAAAAAAAAAAAAATACAAGATCTGAAACTATGAAACTCATAGAGAAAAACAG GAGAAAAGTTTTATACCATTGGTTTTGGCAATAATTTCTTGTATACGACACCAAAGAACA GGCAGTAAAAGCAACAAAAATAGATAAGTGGAACTACATAAAATTAAAAACTGATGCAC AGAAAATAAATAAAAGAAAAAACAGAGTGTAAAAGCAAACCATGAAATGGGAGAGAATA TTTGCAAACCATATATCTGATAATGGGTTAGTATTCAAAATATATAAGGAACACCTACAA CTCAATAGCAAAAACTAACCCAATTAAAAATGGACAATGGACCTGATGGATATCTCTCC AAAGAAGATGTAAAAACAGCCAACAGATACATGAAGAGTGCTTAACATCATTAGTAATTA GGGAAATGCAAACCACATGAGCTATCATCTTACACCTGGTAGGATGACCATTATG AAACAAAAGAAAGAATTAAAAAAAAAAAGTGTTGAAAGGGATGTGGAGAAACTAGAA CCTTTGTACAGCCACTGTGAAAAATGTTTGGAAGTTCCTCAAAAAAATTAAAAATAAAA CTATACGATCCAGTAATCCCACTTTTAGATACTTTTCCAAAATATTTGAAAACAGGAACT CAAAGAGATATTTGCACTCTCATGTTTATTGTAGCCTTATTTACAATAGTCAAGAGGTGG AAACAAATGAAATATATAATGACAGATGAGTCAATAAAATGTGGCATGTACATATCATGG AATATTATTCAGCATTACAAAAGAAGAAAATCTTATAATATGCTGCAACATAGACAAACC TTGAGGACCTTATACTAAATAAAATAAACCAGTCACAGAATGACAAATACTGCATGAATA TACTTCTATGAAGTATCTAAAGTAGTCAGTCATAGAAGCAGGAAGCAGAACGGCAGCTGC CAGGTCCTGGGAGTAAGAGTAAGAGGAAAGTTGCATTTCAGTGGGTATAGAGTTTAAAGC ATGCAAGATGAAAAAGCTCTAAAGATCTGATGTACAATAATATGCATATAATGAACAATA TTGTACTGTTCACTTAAATATGTGTTAGGTCCATGTTATGTGATTTTTACCACATTTTTT TGAAAGCAAGTTGCTAAAGAATTTGCCAAATGGAATTATAGTGACACGAGTTCAAATAAA ATTAAAAACGAGAAACAGTAGAGTTTACTTAATTTGTTAATATATCCATATTATCATTT TGTTTTAGTTTGTTTCCCATTTTTATGTAGCTAGACTGCCAGTTAATCTCCTAAAATTAT TGGCACCATATTTCCCATTTTTTCTGGCTTTTTTATTAGTAACTGGGATCCTTGCAGCTG TATCTATGTGATGCCAAACAATTAGGTTGATCAATTCTGTGACAACAAGCCATCTGGTTA CTTTAGTGAATAGGCCCTTACTTACCTTTCATAAGTTGATTCTATTCTCCTTTGTGCCTT CTCTTTAAATTACCATTATCCTGTAACCATAAATTAAAAATACAGCATCGCTTTTAAAAC GACCCTAATTAGCATTTAGGAACAAACTACACTTGCAAAATTATTTTCGATTGGTAGAGG GAAGAAAAGGGTCTTTTTATTACTATGTATTTGTAATTACTTTTTGTCACTTATGTTATTC TTGTGTCTAAATTCAACTCTAGATTTATTCTCTGTTGATATTTTTTTATCACTTGAGAATA TTTTAGTTTTCAACCTCTATATGGCGGGCTATCACTCCAAATTTAGGTTAAACTGTAGG AGTAGTTATATTTCAGTACTTCTTTTACCTAATCAGCCATTTTAAAATAATTTTGTTCAT GGCAGCAATCTGCATGACAAATTTCTACTTAATAAGCAATGAAATAGTTGGATAAATGTG TATTTCTACATGGGTGAATTTCCCAAAATTCACACTTCAAAGACAGTTGCTGACATTTTT TCAATGAGAGATTTTATTAGATAATGAGTCATCTTAGAGTTATCTTGTAAGTATTCTTTA TGTTTTATAAACATTAGAAATTAAATAGGACTACCATATGGTCTAGCAATCACACTTCTG GGTATATATCCAAAGAAAATCAGTTCAGTATGTCAAAGAGATGTTTCGTATTCATTGCAG

FIG. 1V

WO 01/30991

CTTTATTCACAATAGCCAAGATATAGAATCAATCTAAGTGCCCATCAATGGATAAACGTA GAAAACATGGGCTGGGTGGCTCACGCCTGTAATCGCAGCACTTTGGGAGGCCGAG GCGGGCAGATCACGAGATCAGGAGATCCAGACCATCCTGGCTAACACGGTGAAACCCCAT CTCCACTAAAAAAAATACAAAAAAATTAGCCGGGCATGGTGGTGGGCGCCTGTAGTCC CAGCTACCCGGGAGGCTGAGGCAGGAGAATGGCGTGAACCCGGGAGGCGGAGCTTGCAGT GAGCCGAGGTTGTGCCACTGAACTCCAGCCTGGGCTACAGAACGAGACTCCGTCTCAGTT AAAAAAAAAAAAGGAAAGAAAACGTGGTATATATACACAATGGAATACTATTTAGCCTT TTAAAAGAAGGAAACCCTGTCATTTGCAACAACATGGATGAACCTGAAAAACATGTTAAG AGGAACAAGTCAGGCACAAATACTTAATGATCTCGCTTATATGTGAAATCTAAAAAAGTT GACTTCATGGAAATATAGAGTAGAATGGTGATTATCGGGTGCTGGGAGTTGGGGTAAGAT GTGGTTGGGGAAACGGTCAAAGAATAAAAAATTTCAGTTAAAGAGGAAGAATACATTCAA GAGATCTATTGTACATGTTGAATATAGTTAGTAACAATATTTTGTATCCTCAAATTGCTA AGAGAGTAGATTTTAAGTGTTTTTGACACAAAAACTGATAATTATGTGAGGTAATACATT TTTTAATTAGCTCCCTTTAGCCATTCCACAATGTATACATCTTTTAAAACATCATGTTGT ACATGACAAATATATACAATTTTTATTTGTCAACTTAAAAAATATTAAAGATTTAATGTA GATAAATGAAAGAAAATTAGGAATTAAGGTACAAAAATTATTATAGTGTTTATTATTGG TCTATGTTTACATAGTATTTCTTTGTCTCCATTAGTGTGTTATACAAATACCCAACTAGA AACATGACTTTACAAATGGTGTATCTGATCTTTTATGTCCCTAGTTATTATTTTAGCCCT GTCTTTTTTTTAATAAAACATATTCTGCTTTTTCTTGTCCTCATCCTTCTATGAGTTGA ATTAGTGACTCTACTCCAAAGTAATGGTGTTGCTTTCTCAGACCATATGGTGATACAAAG GCATATGAGTTATCATAAGCATGGTCTGTGTAGGCAAAGCATGTAACTCCACAAATGCTT CTTGAGAGATTCTAATATATCTGTGCCAGACCTGCACAAGGCATAGAGAATAAAAATTT GCACCCCACACAGTCACTCCTCATTCATTCATCAACAATAATCAAGTACCTGGTAATGC TAATGCAGTGTACTATAATTCCATATACATAAACTAATATTTTTAAGATACATGAAGGTT ATGTTATAACTAATAGTCAATGTATTTTTAAAATTACTGTAATCAAATTGTAATTGTAAT TAAGTATTTTCTTAATCAACAGAAACTAAAAGTATAATTTCCATCAACTCCTTTTAAGTA TAAATGTAATTAAATGCCTGGCACATTCTTCACATTATATAAGGATCTTTATACTTAAGA CATTTGGGAAACCCTACTTAGGCTTATCATTGACAAAACATTTTCAAAATCTTTTCATTT GGTCCTCACCACAATACTGTTAAAAAGACAGCCTAAGCTGTTTTGTGCTTCCTCCCTAGT TGGGCATCCCTGTGCAATGAGAGGGACAAACAAGGTGGTTTTAAGGTCAGAAACATCCAA TTGCAGCATCATTGGGAAATTTGTAAGAGCAGCTTTTATAAAATGTCACCAACTCATGTA TCTTTAAAAGATGTGCTGAATCTTATGCCTTGAGATTTTTCTTAGTTTTCCTTATTTTCTA TTCCCCTCCCACTTTCTCTTTGTCCCTTGGTGGCTTCATTAATCCCATATTACAATACAA AGTAAATAATAGTGCTCTGAAGTGCTTCCTATTTGTTCAGGATGAAGTCTGAAAAATGAA ACTGCAATTTTTTTTTTTTGAGACAAAGTCTCACTCTGTTGCCCAGGCTGGAGTGCAAT GGTACCATTTCAGCTCACTGCAACCTCCGACTCCCAAGTTCAAGTGATTCTCCTGCCTCA TCCTCCCCAGTACCTGGGATTACAGGCATGCACCACCACGCCTGGCTAATTTTTGTATTT TTAGTAGAGATGGGGTTTCACCATGTTGGCCAGGGTGGTCTCGAGCTCCTAACCTCAGAT GATCTGCACACCTTGGCCTCCCAAAGTGCTGGGATTACAGGTGTGAGCCACTGAGCCCTG CCAAAAACTGCAATTTTATCTTAGGGGACAGGTAAGCATAAAAACATCCAAAATCATGTA TTTTTTTTTGAGACAGAGTCTGGCTCTGTCGCCCAGGCCTGGAGTGCAGTGGTGAGATCT CGGCTCACTGAAAGCTCCGCCTCCCGGGTTCACACCATTCTCCTGCCTCAGCCTCCCGAG TAGCTGGGACTACAGGTGCCCGCCACCACGCCCGGCTAATTGTGATTCTTTACATTATCA AAGAATTCATGAAAACAGGATATGAAGATTAGTGAAGGATTCTTTTCATTAGCAAAGTAA CTTTTCTTATTTCAAATTTAACACATCTATTTATAAAAGTTATAGAATTTAAAATTTTAAA ATATGAATGAAGAAAAACAAAATCAGCATAACATAGTAATACATATAATTGATATGTACT ATGTGTTACAGTTAGGGCTTAGAAAGATTTAAGCACCTAGCCAAAATTATGCATTATGTT AAGTGGTTGATATCCACTTATTGACAAATATGTATTGAGTATTGAGAATTAGTCATGGAGATA TCAATGGGTTATTTTGATTACTTTTTCCATTACTCCCAAGTGGTCAGGATTAGTTTTAGA TTATTTAAGTAGGTTGGCTGAGTTCACAAAAGCTATTACTATGGGGACCTTAATTGAAAT GTGTGTTTGCACACATAAAACCTGTTCTAATTTTATGCAACATGGAAAGCATTAATGTT TAACATGTATGTTTGAACAGGGAATTTTGTACTGCATTAAAGATTATTCCTGTGTATTAC ATACAATCAAATATTTGACTATTGACTGTCTTAGTATGTTCATCTAATTGTTTCCTATTC CCATGAAAACTGTATCAGTCTGAGAACAGCTACTATATGATATGCATCACTAGTCTCCCC

FIG. 1W

ATGGTGCATAATACTTGATATAAATTAGATGCTGTTGGTTATACTTGGCGGGGGGAAAGG GGACACTAAAAAGGAAGAGTCAATTTCTACTGTGAACAAAGCAAAAAGCAAAAGGAGAGA TAAATGGAATTAAAATTAAAAATGAAATTGAGAGTGTAGATAAATCTATGTAATGAAGATG CTAGTAACATAGGAAGAGAAATAAGATAGGGTATAACAGTGATTATTTTTCCTAATAAGT AGTGTCATGGCAGTTGGAAGACAAGAGATTATCCAAGCACTGGTTATAGTCTGAAAGATG AGTCAATTAAGACTTATACAAATGCAGAAGTCATGGTTGAGGTAGTGAGAGGATTTCCAG GACAGTGATGAATAACAGAACCTCAGCAGAAGGAGCATGTGGACCCAAAGCATCATACGA ATAATGATAGGACCAAGGGAAAAGAAGTCAAGCGGAATGGGGATAGACAAAAGTTTTGAA ATTTATGTGTAAGAGTTGAATGAAGAAGTTATTAATAAGACTTACACAACAAAGAATTT CTACATAGAAGTTGAAAAGACAGCAACAGAGTTTAGAGTTTAGGAAAAAAATTAAATAT TAAATTTTAATATGTAATATTGTAGGATTTGAATACCTTAAAGCTGAAATTCAGTTTTTG ATGCTGCTTCTTAGCATCTTTGTCTTGACATGTATATCAAAATGTAAGAATGTCTGTATC ATATCCAATAACATGCCCCAATGTTTCACAGGTATCACACCAATAGCCCCTGAGATATTG TCACATTCCATTTATCTGCAGAAGTCTTATTCAACTTTCTGTATTAAGTACCAAGAAATT TCTTAGGCAATTAGTAAGTTCACTTGTATTCTTAAAACTTCACAGAATGAAAAATTAAAA ATTTTAATCTCTTTTTCTAGAACAATTGTTTTACAAAGACTTTTCAAGGTTTTTTAATCC TATTTTTTGACAAATAACATATTTTAATGAAAGTAAACATGTAGAAATGACTTAACCAA AACTAGCTATTGACAACTTTTCAGCACTTTTTTTTTGGGTGAATTCAGGAACAAACTTTGT ATTCATTTTATTAATCCACTAAGTAGGGTTGCTTCACTTCCTTGGTTACTGTGCATGTGG ACGAGGCTGATTTTCATGGTGGGATGTTAAAAGGAGGGATTTTTGCAAATCAAACCACAG AACCATCACCTCACACTTGTTAGGATAACAAACATTAGCAAAACCAAAGATGACAAATGC TAGCAAGGATGTGGAGAAATTGGAACTCCTGTATATGCTGACAGAAATATAAAATGATGC AGCCACTATAAAAATTTTTTGTTTTTGAGAATGTGTCTTGCTATGTTGTCCAAGCTGGCA TCAAACTCCAAGACTCAAGTGATCCTTTCACCTCAGCCTCCTGAAGAGCTGGAACTATAG GCATGAACCACTGTGCTGGCTTGGAATATTTTTTTTTTCCTCAAAAAATCAAAAATAGAA TCACCATATGAGCCAGCAATTCCATTTTTGGGTATATATCCAAAATAATTTAAATCAAAA TGTTGAAGAGATATCTGCACTCTCACATTCATTGCAGTAGTCTTCACAAAACAACCTAAA TGTCCATCCATGGATTAATGGGTAAAGAAAATATGGTCTACACATACAATGGAATATTAT TCAGCCTTAAAAAAGAAGGGTATCTTTCTGAATGCAACATCATAGATGAACCTGCAGGAC GTTATGCTAGGTGGAATAAGCCAGGTATAGAAGGACAATTATTGCATGATTCTACTTACA TTAGGTATTTGAAATAGTCAAACTCATGGAAACAGAGACTAGAATGGTAGTTGCCAGGGG CTGGGAGGAGGCAGAAATGAGGAACTGCTGTCCAATGAGTATGTAGTTTGAAATTATGAAA AAATGAATAGGTTCTAGAGATCTGCTGTACAACATTGTGCCTACAGTTAATGATGCAGTA TTATGCACTTAAACATTTATCAAGAGAGGAGATGCCATGTTGAGTGCTCTTTTCACAATG AAAACTACTTGGGAAACAAAATGGAAGGTCCCCAGAGTCGTGAGGGAAGTAAGGTATGGT TCTCATGCCTTAATTATAGTTTACTGCTGAGATAATTGAGAATGAGAGCTCATATTTACT AACCAGGATATGAATAGACTGAGAACTTTAAATAACTTTCCTTTAATTCCATAAAAATCT CCATTCTGTTTTAAAGTCTTTAGTACAGATTTTAGATGTAATAAACTGCTAAGATTTGAG CAACAACTATAAGCATAATAAATGGTTTGCTTTATGGGCAGTTTTACACTAATGCCTCTA ATAATAATAACAGTAGCAATAACAAAAATGACAGGATTCTTAGGACTTCATTACTCAGAG CATAATCCCTAGAAAGCAGCAGTCATTATCTAACCCAGAAACTCCCAAGAGTTTGCTTAA CACTTTAAAATGTATAATCTAAATTAAAGAAAATATGAGTAAATGGTATTGTTTCCCCTG AATTGAAGTAATATGGGATGTGTTGAAAGAATACATCAAGACATTTTTCACTGTCACCTA GCCTGATGACTGACATAGATTAATTACTACATAAATTTCCTCTTCCATTTAATACTGATA AACAGATTTATGGGACTTAAACCACAGTACACAGTTTTGTATTTTTGTACGAAATGGATAA TCACATTTTAAAACATGTGTAAGGCATATTTGCAAACTTGAAACGTCGTCTTCCATAAAT ATATGCTGAATGAATTAATGAATAAAAATTGAGGCAAAAACTCAGGTGTGGCTCAG ATTTCTAGCCAATAAGACCAAGGTTCATTCACTTCACCTCTGTATAGAATCCTTTGTTGG GGGCTGCGAGGAGGCAGTAAGAAGTATCACATCTAATCTTTTCCATAATTAGCCAAGTTA GTTGGTACTTCCCATAACTCTGATACCCATAGGCCCTTGCTATTTCTAGACTTGAGTGTC ATTCAGAAATATGGTTTAGGCGAGCACTAGGAAAGATACACAGTTTTTCTAAAACACATT

FIG. 1X

25/51

ACATACAGGTATTCTATTAAAGGAACTTTTTAATGTTTGACCAGAAAAAATTTCAATATC CCTTTTTATTAAGTTTAAGTTACTGTAATGAAATTAAACATGTGAAGGGAGACTAATACT CTCTTTTAAGAGAAGTAAGAATGAAATATCCATATAAAATACACTGCATTATTCTCTTTG AGAAAGCCAAGATTTTATAAAGTAAAAATCTGCTTTGTGTGCCTTTCCAAATTAGAAGAG TTGAGTGAAATAAAAAGAAATTGACTTACTTGTTAAAGAGAAAAGATTGCCAAGGCTTGC ACGTGCCTTCCATTTAATAAATGCTGTATCAATCTAGCTGTTTCTCTATTTTTAATCATA CATTTTGTTGTTGCTCTAAATTTAATCTTACCTTATACATTGTATAATAGATGTCCCTTA CACTCCTCCTGCTGTGATCAGTCTCTCCGTTTTTGTTCATTGTCCATCATCTTCTACAGA ACAGATGTGTCCTAACCCACTTTCCTAAACACATTTTTGTATACAAAATAATTTCCTTTT TTTAATTTCAGAACTCTATTCTGACAAACATTTGGCTTCAACCTGTÄATTAAAAACTTAA CAATACTTAATAGTTGCCTCAAAGAGCATCCCCTCTTTGTCAATGTGAGACTATTTACAT TAATTTACATGTAATTCAGTTTCATACTCATTCACTGGGGTGTGAATATTAGTCAAACGG GCAATTAATTAATACAATCTTTATATATTCACTTATTAAAATGCACCACACAATTCCTAA TTTATTGAGAGTTCTCACTAAATCTATGGGATGTAAATTTTGAAACAGCTGCAGCTGTTT TCCTTGAATAGAGTCGAAATTATGAATCTAACTTTCTCCGACATGTTGTCTAAAAGGATA TCATCTTACCTTACTCAGTGTGAGCCCTAAAACTAGGAAATGTTTATCAATCTCTGATTG CAGATCAAGTTTAACTATCAAATACAGATTAACTTTTCAGCAAAAATTTGTTAAATATTC AGAGATAGAAATCTTGATGTTGGATGACAAAGATCACTTGTGAAGAACTTTATTAAGTTT TATTTGGTTGAAAAATCTATAATTTTTAGTGAACAACTATCATCCATTATGTTCCAAGCT TTGTGACAACTGTTTTTATGTCCATTAAAACAGTCCTATAAAATAGGTACAAGTATCTCA **ATCTTATACATGTCAAAACTAAAGCACAGAGATGCTAAATAACTTGACTAAACAAGATAT** TGAAGGTGAAGTCTGAGATAGATTTTTAACTCCGAAGTGCATAAACTTTACCTCTATATT ATCTGTCTTCAAAAAGAATGATTTTAAAGATTAGGCTTTTTTATTTCAGAAGAAAATATT AAAGTGAGGTACATTTTAAAGGAAAAGTGACAGACAAAAAATGGATTTTTGAAAAATGAA TAAAGCTGCTTTTTTTTTTTTTGATGGTGTCTTGCTCTGTTGCTCACGCTGGAGTGCAATG GTGCAATCTCAGCTCACTGCAATCTCCGCCTCTCGGATTCTAGTGATTCTCCTGCCTCGG CATCCCGAGTAGCTGGGATTACAGGCGCCCACCACCAGACTCAGCTAATTTTCTGTATTT TTTAGTAAACATGGGGTTTTACCATGTTGGCCAGGCTGGTCTCAAACTCCTGACCTCAGG TGATCCACCCACCTCGGCTTCCCAAAGTGCTGGGATTACCGGCATGAGCCACCACGCATG GCCAAAGCTGGTTTTTAAAAGGGATCATTGTACATTATTATCAAATTTCATTTGAACGTC AAAAATTCTGAGGCAAGAAGGAAATTGAGCCCAGGAGTTTGAGACCAGCCTGGACAAAAT GGCAAGACCCCATCTTTACAAAAACAAAAATAAAATAACACTAGCCAGGCATGGTGGTGC ACACCTATAGTTGTAGCTACTTGGGAAGCTGAGGTGGAAGGATTACTTGAGTACAGAGAA GAGGTTACAATGAGGGAGGATCGTGCCACTGCACTCTAGCCTGGGCAAAAGAGCAAGACC CTGTCTCTAAAGAATAACAAATAAATAAATAAAGTCTGGACAAGCCTAAAATCAGTAATA TTTGGGGAATATGCAAATAGTCTTTGCTTTATTTACTCAATTATTGAAACTATATTCAAA AATAGGAAGTAAAACATGATTTAATATTTTTAGTAAGTTAAACATGTTATAATAATTTG GAAATCCATGTATGTTAGTTAAATATACATTACTATAAAATGTAAAATCAGTGTGGTTTGT AGCAGAGACCTGGATTTTTTATCTTTGTAGTGTACCTACACCATCACAGAAAGGTTTGCC CATTCTGTTGTTTTCAGCCTTCATCTAAGACACTCTCAGATACTATTTCAGGAATTTATG ACAGCAAAATGATATAAGGTGACAAAGTAGAAATAGGTGCTATGCTGCTTTACCTATATT GAGTTATTTTCTTCTCCCAGGATCAGATATTAATGATAAATTCTCTAACATCAAAAAAT TCTACTATGTTTAGAATGGAACACCTGACTTATAGAAAAAAAGGTAAAAGATTGGTTTTG ATTTTAAAATACGATAAGAAGAGGGGGAGAAATGGCTAGATTAATTTGAGGATTACCTA GTGTTAAAATAAGTCCAGATTTAAATCAAGTTTATTAATTCTGAAAAAGATCACATCCTA

FIG. 1Y

26/51

AAGAAGGCATCAAATTGACCCATAAATGTGGATAAAACTTCTGTAAGATAATGAAAGCCC TAGAGAGTAATGTTCAACTCCATTTTCTAATTGGCAACAAATGTATAATATGGGTACACC AGAATATCTAACTCAAAAAGTGGGGGAAAAAAACTCAAAAAGTACGAAATGTTGGCAAAAA TGCAGACAGCTAGGACACTCATACCAGCTGGTAAGTGTAAAAACTAGTACTGCACCAAGC ACTTTAGAAAACTTAACGGCAGTTATGTAGTAATGGTGATCATATGCATACTCTATGATA GCAATTTCACTGTTAGATATATAACTAACAGAAATTTGCACATATGTGTCAGAAGACGTA CATAAGAATGTTAGTAACAGCCCTGTTTACAATAGCCCTGAATTAGAATGAACCAAAATT TCCATCAATTGTAGAGTATTTCAATGATAATATAATCACACACTGGAATGAAAATGATGG ACACAAAATAACACATAAATGTTGATTCCACATAGATAAAGTTAAAAACAGATAACAATT AATCTATGGTGTTACAAATCAGTATACGGATTTCCTTTTGTTGGCAGGGGGGATGTTGTT GGAGAGGAAATAGGAAGAGAGCTTCTGGGGTGCCGGTCATATTGTACTTCTCAGTCTGAA TAGTAGTTACAAGGGTATGTACACTCTGCTGTAATTTGTCCAGTGATACATGATGGTTTG TACATTTTTATACATGTGTGATAATTCAATAAAAATATCTGAAAAGCTACAACAGCAGTG GCAACAACAAGCCCATTAACCACAAGAAATAATCATGTAAATTGTTTTCTTCAAATAAA TGTGTTGTAAATAACTTCTCTCACTCTTTTGGCATATATTTTTGTCCTCTTTTGATATACC CTAATTTTAGGTTTGTTTAATTTTTCAAACATGTCCTTTATGTTTAATACATTTGAGGAA ATCTGCTTAAGAAATGCTTATCTACTCCAACATCTTATCAATGGGAATTTTATTTTTTTA **ACTGTCAAATTTAGATCTATAAGTAACCTGGAATTTATGTTTGTATATGATGTGATGTAG** AAATCAAATTTTTTTTTTTTTTTTTTGTAGATATCAATTTATTCAGTATCATTTGTAGAAAA GATACTTCTTTGATAATGCAGTACATGGCACTTTTGTCATATGTCAAGAGTCCTTATATA ATTCTATTCCCTTGATCTACATTTATATCCTTGTACCAGTACTATACTGTTTTGTTTACT GACACTTTGTATTAATATTTGATAGCTAATGTAAATCCTTCAAATTTGTTTTTCCATAAT ATAATACTGACTAATTTTGGCCCATTATATTTTTATATAAATTTTGAAATCAGCTTGCCA GTCTTTACCAAAGGAAAGCTAGCATTTTAATTTGGAATGCATTGAATCCATATATCAATT TTAGAGAAAACTCACAGCCTTACAATACTTATTCTTCGATTCCATGAGTAGGGTATATCC CCCTATCCATTTAGGTTATTTTTCATATTCCTCATATTTTACAGTGCAGAAATCATGTGT TTCTCATTATTTTTTCCTAGATGTTGAACATTTATTATTCTATTGTCAATAGTATCATC TATTTAAATTGCATTTTCTAGTTGTTTTTTTTTTAATAGAAACATAATTGATTTTGCATAT ATACATTATATTTTATATATCAATTTCATGTGCTCTTATGTTACATATTGTTTTATATTC AGCAAGTGTTACTAAGGTATTTATTAAGATTAGTAGTTTATCTGGAGATTCTTTCACACT ATTTTATTTATTTTATTGCTTTATTACCCTTGCTCCAGCACAATGCTAAATAGAAATTA CCATAAAAGACTTTGTGCACTTACTCCTGATCACTGAGGGAAAGACTATTTATGTGAATT AGTATTTGTAGATATTAACTTTTGAGAATTTAGCTGTCAATCCCAATATGACAACTTGGA GGTGATGCATTTTTTCTTCCTGCTTTAAGATTTTTCCTTTCGTCACTGGTTTTTCAGCA GTTTTATGATAATATAAGTGGGTGTGATTTTCTCTTATATTTATCCTGGTTGAAATTTAT AGCACTTCTTATATCTACAAATATATACCTTTAATTCGTTTTGAAAAATTCTTAGATAAT GTATTTGCCTTGCCAATATCTTTTTAAAGATTGCTTTTTGTCTCATGCTACTTCTATACAC ACATATTGAGAATCCAATCACAGGTATAATAGAATTTTCACCATGTGTTATGCACACTCT TCTGCATTTTCCTTTTTTCCTCTCTTTTTAGCTTGGATATTTTCTATTAGTTTGTAT AATCCTATTAGATGGTTTTATCTAATCTTTCTTTTCTGTTAAATCTCTTTGTTGTTGTTTTCC AGTTCACATATTTTTAAGTTCTATAATTTCCTTGGACTATTTTTCTATTTTTTATATTCT TTATAATATATCTACTTTCTTGACATTATTAATTCAATCATTTTAAAATTTCTGAAATAT TTTATGAAAAATTGTAGAAATTATTTTATGTTCTAGATAATATTATCTTCTTTCACAGAG AATTTGCTTTTGCCTTTGGCCAGCAGCTAGTGTTGGGACAGAAACCACTATCCCGTCAGT CACTGGAGGCTTTGGAAGCTGGGCTTCATTCTTTAGGAGAGCTTGTCTACTTCAGATTTA TCCCTATCAGAGTTCATAACTTGGAGTTACAGCTGAAAGCCAGGGTTGTTTACCTACTTG ATAGGCCTTGAACTCCAATTATCATCTTATTTTTGGTTAGGTACTAAATTTCCGGCTCAG CATCTCATATTATCAGCTTTGTTCTCTGTTTCTCTCTCTGTTCTTAGCTAGAGTTTGC AAATTGCCAAAAACTTTGAGAAGAAAAGAGGCTAAATGCCAGAGCATCTCCCTCTTGCAT TTTCTCCAGGATATTGGCCTTTGATGTCCCTTCTGCCTTAGTAGCTTTCCAATGTCTTAA AGAAATGTGTAACACTTCTGGTTGTTTTAGGTGGGAAGTTTGTTCTGCAGTAAGCTTATC TGCCGTTACCAGAAATAGAAACTATTTTGTAATAGTAAAACAAATGTATACTTTCGTACT ACAATATTTAGTACTTCAGAGAACAATTGGCACTTTCTGGATATTCTCAACCAGGAGTAT GTGGTTGAAACTGCACAGTTTTCTGGAGATGATTTAGGTTCTTCCCTTCTTACTCTAATT

FIG. 1Z

CTGTCACTGGTTGATCTTATCCACTCCACAAGCTTTAATCACAATTTCTATTCTGATGAA TCCCAAATATTTACATGTAAAGAAATTATATCCCCTGGAGTATAGAACCATAAATCTAAA TGCCAACTGGGTATTGACACTAGGATAACTCACAGGTGCTTCAAAATTACATATACAAAG TTGAATTTCTCATCTTCTATCTACTCTTACAAAGCTACCTCATTATCCTTTATCCCCTAG CTCAGTGAGCATCCCCAGCTGTCAAGCAATATACCTGCTAATCATCCTCAGTTCTTCTTA CTCTCTCATCCTCATATCTAATCCCTCACTAAGGCCTGATATTTCAACCTCGTTATTATT TTTGGCATTCACCTTTTTCCATTTTTTGGTTACCAACTTGCTTTCTTGGAATTTTAAAAC TGTCAGTATTAATCTCTCTGCTTGCAACATAAAGACATATATTTTCCACATATTCCGCCT AAGTAATCTTTGAAAAATAGTAGTAAGATATTGCCATTCTGTTGCTTAAAATCTGTCAGT AATTTTGTAATTTTCCAATTCTCCATAGTCTGTAGGACAATATCCAAATGTTTTAACTGA ATACACACACAGAAACACACACACACCACACACACATTTTATGATTCATACTTTGAG TTTAATTGAAAGATAGAACATCTATAAGATGAAAACAGTTGTAGTCAGAGATTCTGGTAT GCAAAGTAGGAGAGAGCCAAGAACTAGAGGTATAACTTTGAATTATAATATTGGGTTG TGAAAGAACAAAAGACAAGTGGATTAAAGACAGATATGCGGTGAAAGAGAAAAGCATTTC TACAGAAAAGACCCCCAAAATAAGTTCATTGCAGGTAGTAAGATGAACAGAAGTCAAATG AAGGACTGACTTTATGGCCCTGTAGAACTCTGAGAACAGGGTCAAAATCCAGATGCATTT CTAAGACATCACACTGGGAACGGGGACTTGTAATGAGTTATCTACAAAGTGTAAAAAGAT GTGGGTAACCAAAAGGTTGTCATTTCCTCCAAAACAAATTTTCCTGGAGTGAAACTGTAA CTACCAGGTATAGTCATTAATAGAACTGCAGACACTAAGACTATGGAACCTTCCGTCTTC CTAACCTTCTCCTCAGGCCAGCCTTAAAGGCCTGTGAAGATCTATTAATAACACTGCTGT TTTGTTCTCTGGCAGCTCTTGGTGCCAGAAGGCTTGGTGCCAATTTGTGGTTGAGCCCCT CCTTGGGAGAAATCATGCCATTCAGAGACAGCTGATAAGTCAAGCCTATTTTCCCACTTT CTTCACTGTATTTTTCCTGTCTGAAGAACTTGTTTATGGATTTGATTTCTGTAGAGATAA TAATCACAGGATTCAGTGGTATAGCATTCCTCTATGCATTTTCTCCCTGCACATTTGTGT GTGTGAAGATACTCTTTCTAAATCCCTTTCAAGACAAATTATTAATTGTGATATATTAAT TATTCTCCACTGTACCTAACGGTTATCAACACTACAGAGGCACCATTGGTTGACAAAAGT GAGAGCTTTTCTCAACATTAACATAATGAGCAAGTGGCAATGAGAAAATATTTGTCCAAT TAGAGACTTTTATATTTTCTTTCTTGAGGAAATAAAACCCGAAACACATTTAAGATACA CACTCTGTCGCCCAGGCTGGAGCACAGTGGCACGATCTCGGCTCACTGCAACCCCCGCCT CCCGGGTTCAAGCGATTCTCCCGCCTTAGCCTCCGGAGTAGCTGGGATTACAGGCGCATA CCACCATGCCCAGCTAATTTTTGTATTTTTGTAGAGATGGGGTTTCGCCATGTTGGCCAG GCCGGTCTTGAACACCTGACCGCGGGTGATCCCCCCGCCTCGTTCTCCCAAAGTGCCGGG AGCACTTTGGGAGGCCGAGGCAGGGGGATCACGAGGTCAGAAGATCTAGACCATCCTGGC TAACACCGTGAAACCCCGTCTCTACTAAAAATACAAAAAATTAGCCAGGCGCGGTGGCG GGCGCCTGTAGTTCCAGCTACTCAGGAGGCTGAGGCAGGAGAATGGCGTGAACCCGGGAG GCGGAGCTTGCAGTGAGCCGAGATAGTGCCACTGCAGTTCGGCCTGGACGAAAGAGCGAG AGATAACCATTTGGGTGGCACATTTCACAACACAGATGCACTTCTTAAGAGTCCTCCATC CGTCAGCGTTGTAAAAAAGGAAGTGGCACGTTTGCATGTAGTTCTTCTGAGACGGAGATT TAGGGACAACTTTGCCAAGGTGTGTAGGTGGAGAATGGGAGATTGAGACAGGCATATTGG CTCAGGAAGACAAGGGAGTAAAACTAGCAATAGAAAGGAGGGCCAATGCCGTAACAGTGT CTTTTTAAGATAATACTCCTGGTCAGCTTCCCAGGTTCTTAAGTCTGGATACTGTAATGA GTTTTATTTCACGTTCTCTTATTTATATTTAATTGAGATGGTGTTGGCCATTTTATCCTT TACATGTGCACAATGTGCAGGTTAGTTACATATGTATACATGTGCCATGCTGGTGTGCTG CCCACCCCACAACAGTCCCCAGAGTGTGATGTTCCCCTTCCTGTGTCCATGTGTTCTCAT TGTTCAATTCCCACCTATGAGTGAGAATATGCGGTGTTTGGTTTTTTGTTCTTGCGATAG TAAATATACTCAGTTCTACATTATAAAAAGTATTACAATGAATTTAATGCTTAAAACTCA

FIG. 1AA

28/51

TTCCGGAAGTGACGATGGAAGCAGGTTCAAATGCTTTCACTGACACTTTGTGGCAAAGTG TGGAACTACAGTATATTTTTCCAAGTTGTTTCCTGATATATTTTTTTATGTACATAACAAT CAATAAATTGTTATGCTATTTATTTATGTACTTATATGTAAATTAAACAACCAAGAAATC GCAAAGTGTTTTATTAAGATGATATCTAAACTGAAATATCACAACTTACTACAAATAATA CTTTGTTTCAAAAATAATTTGAATTGCATATAAAAATCACAGTTGCTGATTAACATTG AATACCATATTTATAAAATGAGTCATTAAGATTATCCCTAGGCATTTTCATTCTGTATTG TCGCCCAGGCTGGAGTGCAGTGGTGAAATCTCGGCTCACTGCAGCCTCCACCTCCCGGGT TCAAGCGATTCTCCTGCCTCAGCCTCCTGAGTAGCTGGGATTACAGGCTCCCACCAA GCCCAGCTAATTTTTGTATTTTTTAGTAGAGACAGGATTTCACTATGTTGGCCAGTCTGG TCTCGAACTCCTGATCTCAGGGGATCCACCCACCTCGGCCTCCCAAAGTGCTGGGATTAC AGGCATGAGCCACCACTCCCAGTCGGCAACTAATTTTTAAAATTGTGGTAAAATATACAT AATATACAATTCAACAACTTAATCAGTTTTAAGTGTATAGTTCAATGACATTAAGTATAT TCACCTTATAGTGCAACCATCGTCACTATCCACCTCCAGAACATTTAAAAATTTTTTAAAA CTGAAACTCTTCACTCATGGAACAATAATGCCTCCTTCCCCTCTTCTCCTAGCCCCTGGG CAAAAAAAAATCTACTTTCTATCTGTCTGATATGATTGCTCTGAGTACCTCATATAAGT GGAATCATGTAATCATTGTCCCTCTCTGTTTTTACCTTATTTTAATAATAATCAAAACTAA ATAAATAAGCAAATTCTTAAAATAAAATTGATATATTTAGTACAGATCCTTTTGAGACAC TCAGTGGTCCACTAATTATGTACCATATCCAATCACATCACAATATCATAAATTTTATAG TCAATTATTAGTTGGCATTTCAAGGCCCAAGTATATGTTTAATAAGAGACACAATCTTAC ATATGCAGTTTACATGTTTTAATCTAGTCTTAGCACCAGCATATCACCTTAGTTTACAT TTGTCTAAGTGCAAGTATTGGTTTTGGAATGTAATTTTGCTCATATACAATCTGTAAGAT ACTAAAACAAAAGCTAGTTTATTATAAGTGAAATAATGGCAAAGGCCATTTTAAAAATAT TGTATTATTTTCCCATTTGAAAATCAGTTTAGTCTTTAGCCCACAAAATAACAGGAAAAT AACTTAAATCATAAAAACTATATCTGAATATTATTTAACATATTTTATAAAGATATCCTT CTTTGGATCATGGCTGCAGATGTTTTCATGCAGCTTGAGCCACTTTCCATGTCTTACGGA ACAGTCTGAAAGATAAAGGAAAAAATAATTGATATCTTCTTGGCACCTCTGCATTTCAA AAATACTATTTCAATAAAGTCCATGTTAGAGGTGGAATTCAAGAATTCACTGAATCTGCA TTCTTGCCTTCTGCTATCCTCTTTTGCCCTCATTTGCTCAATTATTCCTCACTCCTGGTT AATGAAGGCAGGCTTTTAAATACAGACTAACCATAAATTGACTTTAATATTGGTGTTTTAA TGGTTATTCACAGAACTGATTTAAAATGTGGTATCAAGTTCAGGTCCTGGGATTTACCAA AGTTCATCAGAGGACACAGTACATGGCGAATTGAGAACCATAGCCTACTTTATGTCTAAG AGAATATTGACAAACAGCTAAGTTCTCTGTGAGCTCTCAGATTTCACTCAAAAGAAATGA AGAAAGTAAATTCTCTGTTTAGACTTTGTGCCTTTTTTCTCCTTTTAAAGAATTTGCTCA TCGGAAAATATACCATACCAATGGCAGCAACATACTATAAGTTTATGAGCAAATCAATTC CATCCATAGTTACTGCAGAATGTATTATAGGCAGTATTTTTGTTGGGAGAAAAGCAGCAG AAACTTAGCAAAGTAAGGGAAAGGAGAAAAAGCAGCTTATAATGATAAAGAGCCTTTGTGC CCGTAGAGAGATAAGAAAAATACAAAAGAAATCCATAATGATCCACAATAATTTTAGAA TGCAATTTATGGCCATGAAGGGTACAACATGTGATTGGGTATCAAAGAAGAAGAAGTCA TGACCAGTTGATTTTGGACAGTTTTGGATACTATTTAATTGGTTAAAAAGCTATTGAAAT GGAGTATCAACCATTTCCAGACAGAGGAATGGCATGAGTGATGGTCTGGGCACGGAATAT GTTTGACACACTGAAATATCAGATTCACTCTGATGCTCTGTGTATTTTACGGGAAACA TTATAAGGGATAAAGGGCAAAAATTCAACAGAAACCCAGTTACTATTGGCCATCTGAGAA TTTTGTACTGTCCAGGAGAAAAGAGAGCTCTCATTGAAATGGAAGAGTTAATACAACAAG ACATTGTGCTTGTCTGTACTCCTATATATTTTATCCATTAAAGGAATTAATGGATTTTAT CCATTTTATGACATTTATTTTTTTTTTTATGACACTTATCCATTAATGACATTAATGGATAAAA CATATAGGAGTACAGACAGGCACAACGCATGGGGAAACTATTAGGAGGTCACTGCAATAC TATACTTCAGAGAAGCACAAAGTCCAGTGATAAGTTTAAGTTGTATAAATTTAGTGTGCT CTCAGGAGAAGGTGATGTTTACTTTGTACTTTTACAACCTTGCACGGGTGAGTGGGTTAC TGAATAAACAAATAAATGTTTGTGTAACACAAATTTAGAGAATGTGCAGTTGTAGATATA TATGTAGTTCTGAATAGTCCATTTAAAGACAGATACTAGGTTTTCTTCCAGGGTTTCTAG

FIG. 1AB

29/51

AGTTTCGGGTCTTACATTTAAGTCTTTAATCCATCTTCAGTTGCTATTTGTATATGGTGA GAGATATGGGTTTAGTTTTGTTCTTCCGCATATGGCTAATCCAATTTTCCCAGCACCATT TATTGAGTAAGGCGTCCTTCCCCAGTGTCTCTTTTTGTTGAGTTTGTTGAAGATAAATTG CCTGTAGGTATGTGGTTTTATTTCTGGGTTTTCTATTATGTTCTATTGATCTATATGTCT ATTTTTATACTATTAATAGTATCATGCTGTTTGGGTTACTATAGGCTTATAGCATAATTT GAAGTCAGTTAATACGATACCCACAGCTTTGTTCATTTTGCTTAAGATTCATTTGACTAT TTGGGCATAGCCACAGTCTTTAAATATTTGAATGGACATAATGTGAAAACCACACTTAAG ATATGTTTAAACGGCACAGTAATATTATCTAACACAAACTCAAAATTCAAATGTATCCAG TATATATATATATATATCTTCCCCAAAAGTGTGCCTTGGCTTTTAAAAAAGCTTACAA TCTGAGTAGATTGCACCTGGAGAAATGATTGCAGGTATGGATAGCTCACTTAGAGCTATT ACTGATAATCTGAAGTGTGTTCAGAATAAAATAACCAGGGTGATGGGGAATGAAAAGCCC ATAAGTTTCACATGATGGATTCTGATTATCTTTAGGCTGGAGAAGCATAGGCTAGGGAAG TGGGCATAGCTGTTGTTAAATACTTGAATGAATGCCTTTTTGATTTGAATTGTGTTT CTCCAAAAATATATGATTAAGTCCTAATGATCATTACTCAGAATGTGACCTTATTTGGAA ATGGGGTCATTGCAGATGTAATTTGATATGGTAAAGTCATATTGCAGTAGGGTGGGCCTT TAATCCAATATGACTGGGATCCTTATGAGATGATGGCCATGTGAAGATAGAAACACAGTA GAATGTCATGCACTGACAAAGGCAGAAATTGGAGTTATACTGCACAAGCTAAAGAGCACC AAAGATTGCCTGAAAACCACAAGAAAATAGGAAGAGACTAAGAAGAACTTTACTACAGCT TTCAGAGACAGGACAGCCCTGCTGACACCTTGATTTGAGAGTTCTAGCTCCAGAACTGTG AGACAATAAGTTTGTATTGTTTTAAGACACCAGGCTTATGGTACTTTTTTACAGCAGCCT ATTAGGATCTTGTAAGCAGAAACGAAGGGAAAACAGAACATGAACAAGAACTTGCTAGTA ATTAAAGCCACTGCAAAATGAACTCAAGGGCTCCAGCAGGTTTTAAATTACCTGGTATTA TAAATGTTCAAGCAGGATGAATCAGAGATGGTGCAGAGGTGATTATTCATGCATCAGATG GAAGGTTAGACTGAATAATCTCCAAGTGAAAAAATTATATGATCCTATCTTAAAGCCCTG GATATAGCTTGGTGGATAACACTTAAATTGAAGACCTAGTACTTAGTTTTACTTTACTT CTGTTAAACAGAGATATTAATGCTATTCACTTCCTAGTGTTATTATGATGAAACTAGTTA CTTTTGAACATATATTTGCTCTACTATTGTCTAGATTGTCTAGATATAATGCATTAAGTC TTCCCACCAGTGCCATTGCTCGTGTCCAAAATACAGAGTTAAAAGATTAGAAATAATTGC ATGTTTTCTAAGAGTCCTGCGCATTTTCCTAGATCCAATATTGTACTATTTGGACAATTT ATTGACCAAGTACCAGAAATATATTTTTTGCCAATTTTCTCATAACAAACTGTGATAA CAAGTGTTTATAGAAAATAAATCACTCAATGGCATAATTTTCAAATAATAAAGACTACAG TTACCCTGATTAAGGTTCACCTGAGTTTTGGATATTACCACGTGAGAGTTAGAGGACAAT GTGAAGTTTTCAAAATTAAATCCTCTGAAATCCAGGTATCTTGTTAAATTGACATCTGTT GGTAGCTGACAGCCAATTTCAGCTTCAGGAACTAGTAAGAACATTTTCCAGCTTATGAAA CTATTAATAAATGTTACATAATTGTCCAAAGAAATCCTCATTCAGTGATTCAAATTTAAC AAAATTAGGTTTTATTTGCCTATGTAAAGATACTAATCCCTGCATTATTTGGGTGCA TGGGTGACAGCTCTGACAGGTTTGTGATGCCCCAGACAAATTCAGTAACTTTCAGTGAAG CAAACCCATGAATAGATGTGATGGCAGCGTGTACACCTATATAATTCCAGAGCTAGTGAT TATGTAACTTTTATATACGTCAGACCGAAGGAAGACAGAGAATGGAGGAACTGGGTGTTC TTTCAGTAAAGAGCAACTGAATGAGACAGTACATCTTTTGAACTGGGGATATACTACAAG GCAATGAGGGAGGCTGGCTATGAAAGTATTGAAAAATATGTTTGATTGCTGGGTGATGTT TAGAGGCCCTAAGGTAATAGAAAGGAGACAAAATTGAGAGTCTGGAACTTATATGTACTT TATTACAGTACTCTCATTTTCACCAAAGAAGGCAACCCATGTGGTGAAAAGACCACAAGC ATTGGAGCTAAAGCCAAGTTATAGCTGTAGTTTTATATTTTTGTGAGCCCATATGTCCTCA GACAAGTTTCGTGAGTTAATTTCCTTTGTCTCAGCTTCCTCTTTTATAAAATGTGGGTGA TATCATTGTCCCTTAAGATTGTTGTCAGCATTAAACAACATAAGTATATGAACCATCTAG CTCGGTATTTGGCAATGGTAGGAGCTGAATAATTGTTAGCTCTTACCTTAAAAAATTATT

FIG. 1AC

TGTTAAAAGTTCCAAATGCAGCGTTCAGGAGAAGATATGGGTCAAGGTCATGGATGAGGC AAACACTACAATTCAATAAAAATTGTTAGTTCTTAATTTATCTTAACTCAGCAACCGTTT CTTGAGACTCTACTACATATTGAGTACTGAGGGAATAGAAAGATGAATCAAAGACCATTT TTGAGAAATGTCATGCAAGGTAAATGAACTGATTTTAAGCATGTACTTAGCATTCACACA GATTGACAGATTCAGTGAAAACACGGCACAGCCTTCAATTATTTTTCTTTTTAAATACAT ATTTGTGGACTTTATAGAAATACTGACAGTGTTTCCTCACCAATACCTATTTTCTTTGTT GAGTGACTATTCCTTTTTCTTTTCAAATTAGTTTGTGTGGCAGTGTGGAAGAACCACCAC ATGAGGACGGTAACCAACTACTTCATAGTCAATCTTTCTCTGGCTGATGTGCTCGTGACC ATCACCTGCCTTCCAGCCACACTGGTCGTGGATATCACTGAGACCTGGTTTTTTTGGACAG TCCCTTTGCAAAGTGATTCCTTATCTACAGGTAATTGTTTTTAATGCTTTTTTGAAGCTA CTAAAAAGAATGTTCAGCCATAGCGATGGCCCTTATGGTAAATTAACTAGTGAGTTGAGA AATATATTTGCCTAAGGCATTGACAAACTGAAGGAAAATAATACTTGAGAATTTCTGGAG AAATAAGTTAAGTTCTGGGTAAAAATTAAGCAATGAACTGCCAAATCATCATTAGATGCT GCACAACATTTTTGCACAACTTTTTTGATTACTAATTTGATTCCAAAAGTTTGATTTTG CACAAACTTTTTTTTTTCCAAATTTGATCCCAAAAGTTTGATTTTGCGCAAACTTTTTTG ATTCCTAATTTCCCCATTGTTAAATAAGAAACTTGAACCAATTAATGATTTAACCAATTA ATGATCTCCCCAAACCAATTATTGATCTTTCTCTTGAACCAATTAATGATCTGCCAGTCC AAGTCATTGAGCATATTTGTTTTTACAAGTGATTTTATTTTATACTGAAGAATTAAGACC TACTTGGTCAAATCAGTGCCATGAACAGGTTTTAGTGTAGATTCTAATTCAAACTACCGG ATTTGGAATCTCCGTTCTGCCATTCACCAATTGTATGCTATCAAGCCAAATAGTTGTAAT TCACTTATTTAAAAGAATAATTTAAATGAGATCTACCTCATATGGTTGCTGTGACCATTT ACTTACATAATTCATATAAATAAGTTGGCACAGTGATTACCCTCTGGAAGAGATGATCTT ATAAAAACAGTATATTCTCAATAAACATCAATTATCAGCATCAGAATCATCATTACTAGG TGTTTTTCTTTCCTTAAGAGTGAAAACAGCTTCTTTTTCTATTTAATTGCCATTTCAGTA ATTAAGAATGAATACTTTCAGAGATTAGTGTTCTGATTGTTATTATAGCTCTAAAATTTT TCCCTTAGGAATTGGACAGCAAATGAAATGGTGACCACTCTCTGCTTGTCTTCCCATAGC TTTCCTGCACCCTCAGTTTTTACGCCATGCAGTCTCCCAGATGGTGCCTATAATATTTTA TACTAGACATTATGCCGATTAGGCTTTTGGAATGAAATGTTGCAAAGAGATATTTAGTTC AATAGTTCCTCAATTACCTCTTATAAAAAGAAGTGAAAAATTTTTAAGGTTAAACATTGT TTATAGAATAGTAAGTGGAAAATACTATAGAAGTTATAAGCTCCATGCATATATTATGTT TAATTATAAAGCTAGTTTGGATCAGCCTGCTGAAAATCATGAATGGATTACAAAACGAAC AGTAGCACATTTTTTTGTGTGTGAGGAAAAACTACATGGGACAATAGAGAAAAATATTCT CATAGAGGAAAAGTTAGTAAGAAATGAATGGCTCTGGTGGTGTTTGCATAGAGGCACTAG GAAAGTAATACATTTCAGATAATTCTAATATTTCATTATCTCTGTGGTACTTCCAGAAAG CCTTTTACCTCTCTTGGTTTCAATAACTACCCAGGAGAATATTTTGAGGATTCTCTTAAG CTGTGAAACAGAAGATTCCCTGGTGGGAAGTGAAGTGATAAGGGCAGGTGCAGTCATGTG CTAATGCACAGCGATAGCTTTCTGCAGAGCAGCATCTCAGAGTTTCCTGTGAGTATTTG TATAATCACTTACAGTCCTCTTTCACACAGCAGAACTATTTAACAAGTCCTACAGTTCAA GGAATATCCTCATCTCTGGAAGGATTCTGTCTGCCTCTCTGCACACAGTGTCCAATCTAA TCAATTCCTTAGCTGCTCCTCTTCTCCATAGAGCAAGGGAAAAAACTACTGGGTAACCAC ATGATGCAAAAGACTAGATCCATTTGTTACCCCATCTAACATTACTTCTTGATGGAAAGG TGTAAATGCACCAAGAGTTGGTGCACAGGTAAAACTAGTATCTCCAAATTCTTCATATT TATTGCCTCATTTTTCATAGAATGTTCCCAAATGCAATGAACAGTGCCAATGGGCAATAA ACATATAATTTAAATTTGAGCAGATTTTCTCCCTAGTTGTGACATTCTGTAACTAATGAC TTATATCCCTGATATGATATTTATGTCTTACTGAATATTTAAAAACATGTTACATCATGC CCAGCCACATTTTAAAGTTATTTGGTTGCATTTTAGATTACTTGGACGTTTATTAATTTG ATGCTGTAACGCATTGGCAAATGCACAAAAATCTCAAAAGTCTCACAAATGTTATAAAGC TTAGCTGAATAATTAAAATGACTCTTTTGTATCTTTAATAATTGCATAACTCCAAGACCA TTAACATGTATTCAGCTATTTGCTGAACAATTATCATGTATTTCACTTCTCTCCAACAA

FIG. 1AD

CAAACAAGCTCCCTCTTTCAGTCATACTTTGAAACCTTTCTACCTATTAGTGCTTATC ATCCAAATCTGTGATTTGGCAAAATTTTCATTTCTCCTTATAGTGAATCTTTAAGATACC TTTGCCGTATCTATTTGCTAGTATAAAACAGTGGACTTCTCTACTAAAGGAAATCCCCAA ACATTATCCTGTGCGAAGGGTGCCCATAGTATAGGTCAAAGACCAAGTACCTGAAGGCAG AAGAAAGTTCCCATTATCTCACTCCACTTCATTCTCAACATTCATAATCCACACTAGATT CATTTCTCAAATGACTTACTATTCAACAAACTTGAGCTAATATCAGAATCCAAATGAAAA AGACACCCAGAAGTGCACTCTTAGAAGTTAAAAGCAACAACAAAACTTTCACTTATAATT ACTTATGATAAAATGCAATTTTACATCACCTCCAAGAAAATCTTATACATTGCACATAAT TGTATATTAATGTGTTAATTGCACAAGCAAATATAGTAGGTCAAACAATGAATATTAGCT CACTGATTGTCAAGGGTTCATTCAATGGATTGGTTCATTCTACTGTTAGATACATCACAC TAGCATATTCCTCCCTTTTCTGTGTGATGAAGGGCAGTGCTCCCTGGGTCACTATTGGCA CTGGATGTCAGTCTTCCAAGTGAACTGATATGAATTGATTATTATGACCTAATGGCATTA GGAAACACTAGAAATGACATTGATATTTGAACCATGCTACATCTATCCCATTTATCCATG TTGATTAAATTAATGGATTATAAATTACTAAGGCTTGATGAACACTTTGTACTTCTAATT GCTAGAGAGGATTGATATCTCTAGCCCAGAAGCTATGAAAAGGCGACTGTGCGAATCT ATACAACCATAGTTCTATTCCCAGGTTAGCAATGGTATTGAGGGGCCCTAGGTGCTTAAC TTATTTGCAGAGAAGGAATGGAGGTTGTAGAGAATAAGGTGATACTGGTTTGAGAAAGAG AGTTGAAGGTACCCTCAGGTAGCACTAAGAAATTTCTAGGAGTCACTAATCAACTTAAGC CCATTCTCATAGAGTCCAGCCCCTTAAAATTACACTTAAAATGAAATTAGCCTCCAATAA TTTAGCAAAGGTTAGGCTTTCACTTGTAATTTCTATGAATATTCTTCTCTGAAAAGCAAT CTGTTCCAATTAAAATATAGAACTTCAGACTCAAGAATGAAAGATAAAACTAATAGTATC ATCATCATTATTATTATTATAATCATAAGAAATAGTAAACACACAGCACTTATATGCCAG CCCTGGAATAGACATTTTCATCTCAACTAACTGTCCATACAATTCCATGGTTAGGTACTA ${\tt TTAATCATCCACATTTTACAGATGAGAAAACTGAGGAATGGAGAGGTTAAATAATCTCCT}$ TAAGATCACTCCATATGTCAGATGGGATTCATGCCCAGAAAACCTGGTTGCAGACTCGAT TCCAGCTATACTCTTCTGCCTCTCCCATAGAGAACAAAAGAATCATACTTGATAAGAAT AGCTAGCTGATCAGAGAGAGAGACAGATAGTTCATCCTGACAGCCCAGAGACTTTCTGC ACTGTTGCACTGGATCTTAGATCTCTTTCACTCATTTGTACCTATAATCAACATATCAAC AAGAAAGGTCCTCATGTAAAAGACAGAGATAACTACCCTTTCCACATATTATGAGATCAA TATAACCAGGACAGAAAAATAGAAGAAGATGACTGGACTATATCTACTGCCTTCAATTAA GGCTCACCACTATTAATGGATTAACAAATATTTGTTTTAAAGACACATGCAAGTATACGT TCACTGCAGCACTATTCACAATAACAAAAACGTGGAATCAACCTAAATGCCCCATCAATGA TAGACTGGATAAAGATAATGTGGTACATATACACCATGGAATACTATGCAACCATAAAAA AGAATGAGATCATGTCCTTTGCAGCAACATGAAAGGTGCTGGAGGCCATTATCCTTAGCA AACAAATGCAGGAACAGAAAAGCAAATACTACATGTTCTCATTTATAAATGGGAGCTAAA TGATGAGAACACATGGACACATAAAGGGGAACAACACGCACTGGGGCCTTTCAGAGGGTA GAGGGTGGGAGAAGGAGGATCAGGAAAAATAACCAGTGGATACTTGGATTAATACCT CTGCTCATGTACCCCTGAAATTAAAATAGAAGTTAAAAACAAAATATTTCTTAAATGCAT AATGGATATCAAATGTTGTATCAGATATTGGGGACACAGTTGTGAAAAAAACAGAAGCAG AATGTAAAAGAAATCAGAAGTCTTTTTAAAGGGAGAGGGGATTCTGAGAGTGATATCAGA ATCAATATTTCATCCAGTATAAGAGAGCACATTGAACATAATTACATTAACTAATAATGT GGATATATGAATTTTTAAAATTTTTTGTTGTTGTTATTTCCTTAAAGTGTCAAGTTAAAG AATGATTTGTGGCATTGTTAATTATATACAAATTTTGACTGGGTGAACTTACCTAGTTTT TGGAATCACATTGACTAGGCTAGCAGTGAGCAAACTGTCATAAGGAGATTCGCATACAAA ATTCTCTTTTAATATGACTCGTAACTTTCCTTGGGTGCTACATGTTGAAAATGCACTGAT GTACAAATAGCCCTTATTATTTGAAAATATGAAATAAGCTACCCATAATTTAAAAATGTT GGGTATGATTTCTTCCAATCAATGGAAAAATTACAGGACAAAATAATTACAGTAATTATT TAAAGAATGCCATATTATAAATTAAGACATTTGGAGTAAAAAAAGATTGCAAAGTTTTCA TCATACCTTTTCATGTTTAACAATAAATTTACATTTAAAAGTATATTTCTAATATTTCAT TTTTGTGATATAATTTCTTTTTAAATAGAAAGCACTTGCATGGATTGTTTATTTTTGGCA

FIG. 1AE

32/51

GCTTTGAATTTGCTTATATGTTGTGACTACCTTTCTCATATAGTAAATATATTAAGAGTA GGTGTCTGTCTCTCACACTGAGCTGTATCGCCTTGGATCGGTGGTATGCAATCTG TCACCCTTTGATGTTTAAGAGCACAGCAAAGCGGGCCCGTAACAGCATTGTCATCATCTG GATTGTCTCCTGCATTATAATGATTCCTCAGGCCATCGTCATGGAGTGCAGCACCGTGTT CCCAGGCTTAGCCAATAAAACCACCCTCTTTACGGTGTGTGATGAGCGCTGGGGTGGTAA GTACCTTATGGCCCATCAACTGACATTTATATTACAGCAGCAAATTGAAAATTGGATTAG CATAGCCATTGTAAAGCTGGGCTTATATATTTTATTGACATTTGTGAATACAGTTTTGCA AGAGCATGAAAACCAACTTGAATTTCAAAACAATTTCACAGAATAACTCTACCTATCTGA ATCCTTTGGAAATGTTATCTATTATTTTCTCATTTTCATATCTTTTTGGATAGGAAATGAA AGGAGATTATTCTACAATTCAGATTTGATTATTTTAGTTTTTCTTAAACTCTTTAAACAA AAAGCAATATGGAATACAAATCCGATTATGTATTCTGGAATGATCCACGATTTATAAGAT GGTTCAACACTGTGTTGTCTAGTGTCAGGGTCCCTAATGGGCTTCAAATACAACTGAATT TTTTCATTTTAAGACCATGTCCTGGATCACATGGTCCTGGGAACATGGCCAGAGTCAGCA TGTGGTTCTCTAAGTCAAATAATCCAAATTTGTTTTCTCTATTCATAATACATTATTGCT ATCTGACAAGTATGATGTGAAATTTAAGCAATCAGGTTTGAAGGCTTTATGTTTCTTTGG ${\tt TTAGAAATTCTTAGAGTCAGTCTGAGGTTTTTGTGTAACAGTGAGAATACTGCTATCAAC}$ ACCTGGTGCTAGCACAAATCTGGGCACAGGAAAGAATGACAGAAAATAAAATAACCCTGC CCAATCTGACCTCTTCTAGGTAAGTATACTAAAAATGGCTGATATTTAGAGAATTCATAT GTTAACATTGTTTTTTTTTAGAAAGATGTATCAAAACAAGCAGTGCACACCAGGGACTGA TTAAGGATAATATTCTTAAATATTGTAATCTTTGAATTTCTGTTATTTCCTACCTTGGTG TTTGTACTAGAACACCGAAAGGAAAAAAAGCCAATCACTGATATATTAGGCATATACTAC AGGATATATCTACAGCAAGATAATATTTAAGAGAGGCTGGGATTATTTCATATATTGTTG CAAGACCTATAATAACTAAAATTTTATAATTTGCTTTATCTATTACCCCAAATATCAAAT TTTTCTCTGTATCATTTTCAGTTTTTTCCAATTTTCCAAATTAATAGTGCAGACAAAAAA $\verb|AAAATCAATGGAAATTTCCAAAATGGTAGGAATATTTATGAAGTGTCTTATGTCCCATTC|$ ATTTAATGCTCAAACACCACCTTGAGAACTTAGTATATGTCAGGCATTGTGCCCACCTGG AGAGAAACAGACTCTGCTTACGGGAGCACACTCTATATAATAAGGCTCAAAGGCCAATAA AAATGAGAGAACAAACTCAGTTCTGGCCATTTGAACAAAAGTTTACAGAGGAACTGCTAA CATTCCAGCAGAACATTAAAGATAAGCAAAAATTCTCCAGACTGAGAAGAGGGAAAAGGA TGTCCAGAAAGCAAGAAAATCCACATCATGGATACTACATTACAAAGCAGAAAGAGTGAA TCAGCACTTGTAGTTTCTGGAACATAGGGGCAGGTAGTGTAAATAATTGAATTTTGAAAC AAGATGGGTTGGGGACTGACATGTCTCTTATACGATCCTTTTACACTGGTTTAT ATTTAGAAAGCCTAAAAAGGTCTTTCTCAGAATCCTGTATTAAACTCGAGACTAAATTTA ACCCTAGAAAGATTATTATTTTTTTCAAGATTATGAAGCAAATAGGTACATTTAAATCT AAAGCTTCCAACTTGTAAGTTGGGATTCCTTAAGTTTTATAGGGATTGCTATTAGATAAA ATATAAAAATATTTTCAATATGTGTCAGCAGTATTTTCTCTAATATTCCGGCAATTAGT TTCACTTATATGTTTATGGGTTGCTTTTATAAGCTTTTCTTTTTTTAATGTTTCCCTGAA TAATCAAGTAACAGTAACCTCCATTAACAAAAAGATTGCAAAGTCATGGATTCCTGTTCA GTTATTATGATTATGTAAATAGACGTATGATTTTTAAATTACCTCTGAGTGGTAAATATA AATACATAAAGCTCATTTCTACTCTGATATTTTATTACATAACTCTAGCATGGACATTTT CATTAAAAAAGGAAACAATTGTTGAATATGTAAAAACCTAAACTTAGCCTTCAGAAGTC ATTTAAGAAAACTATTTGAAGGTGATTTTATAATAGCCTATAATTAAATGCTTGTAAAGA CTAAAATTAAGTATTATTGGACTGAATTGATTAGCTACAAAATCCAACTTAGTAAAAGCT AGGCAGATTTTTATCTTTCTTATTAGATATTTATTATGATGATTTCTACATAGCATGTAA AATCATTGTTCATGTAAACTATTTATAAGTCCATGTTCGACTTATAATGTTAAACCTTTTG TATATGTGTGATTGTCACAACTTTTTAAAAAACCATAGGAAAGTATATTTTACAGTGTCA TCTCTCTAAATTCAAATATTTTTAAAGGCCAACTGTCATTTAGCCTGATTTTTAAAACTA TTGTAAAATATCTTCTATTTGAGATTAATTCATAATCTGTGTTTCTTATCTTTATTCTAA GTTAAATCAATAATGTAGTTATAAAAGTAGAGAGTAGAATCATAATTATCCTACAACCAA AACTTTTGGAAAAGTTCTGTAAATGCTGTTTTTACTCATGGTGCAAAATAACTGAGAACTC

FIG. 1AF

TGTCTAACTAAAAAATTTACCAGCAATATGTAATTATATATGGATAAATGATTTCTAAAA CTAATTATTCATTATTGCCTATTACTTCTTCATAAAAAGAACCATAAGCCATGATTTC TGGCAGACACACACACACACACATATAAATAATGTAAATACTTATTTTAATAACC TTTAAAATATACATTTGTATGTGTTCACTGTTTGCTTCAGTCACATCATTTCATACTTCT AAAATTATTAAATTAACCCACAATTTCTTGCTTGCTTGGTTTGTAAATGCATAATTCTAC AGGAAAGATCCTACAGAAAGAAATTCTTTGCTGGGTGGTGGGCTCAAGCCTGTAATCCC AGCACTTTGGGAGGCCGAGATGGGCGGATCATGAGGTCAGGAGTTGGAGACCAGCCTGGC CAACATGGTGAAACCCCGTCTCTACTGAAAACACAAAAATTAGCTGGGCATGGTGGG CGCCTGTAATTCCAGCTACTCGGGAAGCTGAGGCAGGACAATCGCTTGAAACCGAAAGGC GGAGGTTGCAGTGAGCCGAGATCATGGCACTGCACTCCAGCCTGGGCAAAAGAGCAAGAC TTCACTATTCATCCAGTGTTTAATTAGCATGTACCCTTGGTCAATTGTTCTGGACA CTGGAGATTAGTAGCATCTCTCTTTTTGAATATTACTGACAAATTGTTCTTTGGTAGGCT AAAAAAAAAAATGGAACCATTTTTACAGTCAAAGTAATTATGGCATCTGGCCTATTATG AGGTTTGAAAGCATATAAATATGTGTATAAGTCTATTAATGGGAAGATTTATTAAACATA TTTATTAGGGAGAAGATAGTAAAACATATTAAAGATTCAGGTAAACTTAATGAACCCCTA AACTTTGAAAAGACATTCCATGTTGAATATTGGGAAATTATATTTAATTTACTTGTTCAT TCAATTCCTGATAAGTGTACCATGAAAGAGGAATGTTTCTAGTTTCTAGATAATTAAGAT **AACATGCTGGCTGAATAATGAACCTTAAGTCATCTGAGAGAAATTAAGTTTTGCCTGTCA AATATACAATATAACTCTTTAATCTCTGATTTCAAAGACTAAAGATCCACATTTGTTCCT** TATTAGTTAGTTTCATATATATATATAAAATTTATTTAGATTGTGCTTATTCATCAGT TGAGTAAAAACAGTAATTTTTAATGATTATCAATATTTAAAACTTTTTTAAATTAAAGTA ATGCTTATGTGAAACAAATTTTGTGTAGTTATATTCTAGGTTATATACAAATGTCTTAAA TACATTGAAGACATTGCTTATGAAGTACAGAAAGACTTCAAAGATATTTTCATCACACAT AATTTAAAATTTCAATGGCATATCTGAGTTTTTAATCAGCTTAGACTATCATGTTTCCCT AGTTATCTATTATAATCTCCTTATTCAAACAATCTATCCTACCCTGGAAGGATAATTTTG CTTGATCTTTTTCCATATCAGTGTTCATTATAATAATTTGCATTTAGCAGTCAATTACA TATTTTTTTTAATTATTCATAAATATACCAACCACATAGGAGCTTTTGCTACCATCTATT CAAAACGCCAAACTGTTATCACAGTGATGCTATCCATAGCTGCAGTGGAAAAAATTTACC TCTCAAATCTACTTTCCTCTATCCACTCAATTGGTCTTATGCAGACAACAGGGCTTCGCA GGTATGTAAGCTTCAAAGTTATATAGATTTTGTCATGAGGAAAGCTCATGTGACACCTCT TCAAAACAATAAAAGTTCAAAGCCTCTTAGGTGCCTGGGAAGTGCTGAGATCACTTTCA GATTCCTTTGAAATTGGCCCGCCATATGCTGTGTAGGCTGTGGCACTTCAAAGGGAAAGA CTGTTATTCTCAAGTCAGAATGCTTGAATGTTATCACTTTTTATGTAACTGGCCTGCTT AGAGAGTAGCTAAAAATGACACCTCAAATTGGTCTCTTAGACCTGCCAACACATGCATCC TACTGACCCTGCTGAAGACTGCAGCGGATAAAGACATCTAAACCAAAAGAGAAGATGGGT TTAGAAGCATGAATATGGAGAAAATTAGACTCAAACTCAACTGCATCTGAAAGACAGCCT ATGGAAATAAGATTGTGGAGGATATTAAACTCATAAATATGTTAAAATATATCCAGCAAG AATCAAATGCATGATTGCTCAATAAATATTATCTATTATTATGACAATCATCATGCTTAT TATTGATTAATCCTGACTGTAAACTGCTCTTATCACAAATCTGATCACATAACCAAGCTT TCATGCTTCTACATCCCCTTTATGAAGTAATGAAAAGAATAAAATACATAGAGGTAATAG CATTATTCCTCAACAATACTATGGGATAAACCCCCTTGTCAATAGAAAAGTCAAAACAAA GTATGTAAATTTTAGAAGAAAACAAAACAGCTCTGTTGTGTTAGCATTCAATTAGAATT ATAATGAGTTAATTACATTTAATATCTATGGAATCTATGCAAGATATATTGCTTCCTCTT TTACATTGCAGTAAAAGTAGGTAGACCATTGTGATATATTCGAATACAAGTACAAAAATA TCTTCTAAAATCTACAGGGAACTCAAACAAATCAGGAAGAAAAAATGCAAACAATCTTAT CAAAAGTTGGCTAAGAACATGAATAGACAATTCTCAAAAGAAGGTATACAAATAGCCAAC AAGCATATGAAAAAATGTTCAGTATCACTAAGAATCAGGGAAATGCAAATCAAAACCACA GCCCAGGCTGGAGTGCATGATCTCAGCTCTCTGCGACCTCCACCTCCCAAGTTC AAACGATACTCCTGCCTTGGTCTCCCAAAGTACTGGGATTACAGACGTGAGCCTGTAATT GGTGTCTGGCCAATACCACCTTACTCTTACAAGGATGGCCATAATCAAAAAGCCAAAAAA TTAAGGACATTGGAATGAATGTGGTGGAGAGGGAACACTTTTACACTGCTGGTGGGAATG TAAGCTAGTACGACAACTATGGAAAACAGTGTGGAGATTCCTTAAAGAACTAAAAGTAGA TCTACTATTTGATCCATCAATCTCCCTACTCTGGTAGCTACCCAGAGGAAAATAAGCCAT TATACTAAAAAGATACCTGCACATGCATGTTTACAGCAGCACAATTCGCAAATGGAAAAA

FIG. 1AG

34/51

TAGCATAGAATACTACTTAGCCATAAAAAGGAACGAAATAATGGCATTCCCAGCAACCTG GAGGGATTTGGAGACCATTATTCTAAGTGAAGTAATTCAGGAATGGAAAACCAAACAACA TATGTTCTCACTCATAAGTGGGAGGATGCAAAGGCATAAGAATGATAAAATGGACTTCAG GTACTCAGGGGAAAGTGAGGGAGAGGGGTGAAGGATAAAAGACCACAGATTGGGTAAAG TGTACACTGCATGGGTGATAGATGCACCAAAATCTCAGAAATCACCACTGAAGATTCATG TAACCAAACACCAACTGTTTCCCCAAAACCTATCGGAATAAAAAATTAAAAAAATACATA TATGTACTATATAATATATGAATATATCTAATATTAATATACAATATAAAATTTAATA GAATATATATATATATATATATCTTCTAGAGCATTTACAAAGTTAGTAATCAATATAA TTTAGAAAAGCTAAAATATTAAACCACAATGCCATGAAGTGATTAATCGACTTATTCGTA ACTACAGTCCTAGAGGTGTTTATGCTTAATAAGTGAGAAAATATTCATATTGGATTGGAG AAAATAAATGTTATAAAGCCTTAAAATTCTCATTTTTATTAAAAGTATATACATGTATTT TTAATAAAAGCATACACACCACAGACATACTATGCTTAAAGAGGAATTTTGTATATGT TCCAATAAGTCAACAAAAATAATCATTGTCAAATTTGTATTGTATTTAGTTTTCAAAATT TTTTTCACATTTGTATTTGGAGATACAACTGAGAATAGCCTCCCATTTCTCAGGGAACTT ACATTCTAATAAGGAACAACCAACTGAGTTTATATTTTCTTCCCATTTTAACCAAAGCAT AAATTAATTTTGTTAATTAGCCAGATGTAATCAAGTCAAATAAAGGGCCTTTTAATAACT GAACACTTGACTTTGGGTAGCACAAATTAAGAAATAGCTAATGCTTATTTTTCTGAGTAC ATTAAGTGAAATTACGACTTCACATTTGGCATGTGTATACCCATATACTGAGTAAAATAA ATAAAAGGTAAATATGCACTTTGAAGAAAAGCATTGACATGTATCTTTTTTAAAAGTCCA TCAATTGTAACGTAAGGTTTTGTTTTTTGACTTTCATCCTAGGTGAAATTTATCCCAAG ATGTACCACATCTGTTTCTTGTGACATACATGGCACCACTGTGTCTCATGGTGTTG GCTTATCTGCAAATATTTCGCAAACTCTGGTGTCGACAGGTATATAGTTTCAAATATTTT GTGCTTTTTTTTTAGGATGCACTTATAAACAAAATTTAAGAATGCATTGAACCAATATAA CATGTTCATAAAAGTATTATATTGTGTGTTCTTTTAAAGTAATGAGAACCCAGACATAGA AATATGTCTAGGCATTTTTAGAGTAATATTCAGGAAATGTATTTTATAAACTGATTAAGT ACTTTACATTTTAAATAAAATTTAACATCTGTGATTAATTGTCTTTTGTCTAGGAATAAC ACTAATTTCGCTTTCTATGAGAAATAGCAAATAAAAATTCCTTTAGAGATTTTTGAGACT CTAAGTCTGAAAGGTTATATTTGTAATCAGATTTATTTAAAACATTGGAACATATAGGTT AAATCTCCAACTTCAAAGATCTTATTTTTTAGAATATTATAAGAATCAGGCAGAATGTAT ${\tt AATTTTAAAAACTGTATATAATGCTGATTTGGGGTTACTACACTTTGTTAGATAATTCTG}$ GAACTTTTCACATACATGTCTCACACAAAAGCTAAAAATTCTACTTTTTGCCATTGAGGA ATTCATAGTCTAGAGGGGGGCATCATCAGATGCAGGGCGAAAATTACTTTAAATATAAG CACAGAGAATCAGAGCAAAATGTACTAAAACCATATCTAATACAGGAAAGGTAACATTTA AGAGACAAAAAGGATATGGCTCAGTCTCCCATTTTGTAAAATGTATCTTAAAATGCCA GGCTTGTGGAAGGACATGCCATTAGTCTGTTTTCTCTGACACATTTTATCCAACTGAAAA GATTTACTGGAGTCACCTTAATTCATTAAAAAGATTTCACAAACACTTTATTTGGTCTTT GAGGATGTGTCTTTGTTTTTTAATCAACACTTGTTATTCAAAGCATTTTTCAAGATCAT CTTTCACTGACTGGATATGAGCAACACTCATTTTTTTTAACACTATATGGCTCATAATTT CAATATTTCTCTTTTCCTCTGCTATTACAAAGAAGTCATTTCTTTTATGACCTTACAAG TGAAACCAGTAGCAACATTTATTAACATTTTTGTTTCCCATCATTTTTTACTATAAAAACT AATGTGGACCACTATAAAATATGAGTGGTGATTTTCTAGATGTTGGTGACAGTTTTCTCA

FIG. 1AH

TCCTGCTTCCTAGTACCTATTATTGTATCTGTCAGGTTTTGCTAGGTTATTATTCTTCTA TTAAAAAATGTGGTTTGCAACAACAGTTCTGTTTCACTCCTATTACAGGTCAGTGGGGAG GGCTGGCTGGGCACTGTGCTCCATTTGTTTTCTCATTCCAGAACCTAGTCTGAAGAAAT GGCACTTTCTGGGACATGGCATTCTGAGACTGAGAGAAAAGAAAACTGGAAGAAAAGTA TATTTTCTTTAATGTCTTTTATGAACCGGCATGTGTTACATCTCACTTTTCATTGGCTA AAACAAGTCACGTGGTTAAACTTGATCATGAAGAGGGGACACATTCTTCTCTGACAGAAA ATGATACACTATACAGAAGTCAAAAACACAAGGGCCAGACTGCATGAATTTAAATCCTGA CTCCACCAAGTAGTAGTGACATGAATTTTGTAAATGGCTTAAATTTTTTGTGACTCCCTTT ATTAACTTTAAAATGGGGTTGTATAGCATCTTCCTCATAGGTTTGGTACATGCATTCAGG TGTGTCCAAGGGAGAACACCGTCGTGGGTTCTCAGTTTCTATTTCTATTTGGGCCAGT AAAACCCCTTCCTATCCCTCTTTTCTGCTTATTACTAGAGACAGAAACTAAAAACCAGGG CTTCAGGCTGCTAAAAGCCTAAAACAAAACAAAACTACAACAACAAAATAAGGTG GGTTGGACAAGCTTGCTTAGATGAATTAACTCAAGTGCCTAAATATAGACAGTGCTCATT AAACAAAATATCTTAATGGATGTTGTTTAATAATGGCCTCTCAACTAATTGTACTTACAT TTAAATAGCAAGCATGTGTTGAATTGGTATATGTGACTATTTTTTAAAAAATGCACATTG AAATACCAGTATGGTGCTTCTTATTTGTCTGGTTCTTCTACTCTACTAAGATAAAGATAG TCTCGCTGTCATCTTTGTATCCCTATAAATAGCACGTGCTCAGCACACATCAGTTGCTTT TTTCATAAGAACAAAGTGAGTAGAATAGGAGAAAGTGCTGGGAAAGTTTAGAGAGGACAT AGAGAATCTATTGCCCAGTTACTCCGATAAACATTTGTAGAAATGGATTAGAATCTGAAA AATTTCTTGAAGGGGAAAAAGCAATTAATGAGCATGTAGGAATAAAGATATTTTAGATTT AGATTCAGATTTTGTTGGGGAATGTTCAGTGTTAAGATTATCCCCTATTTCCTTATTTTT ATTGACATACCTTTAAAACTCTTTCAAGGTTGCAATGTATCTGTCTTGTTACTTTTACAT GGTAAAACTTTACCATGATACCATGGTTACCCTAAAGTTTACATGGTACCATCAGAGAAA ATGTTTTAAAAAGTTTGTTAAATGAATGAGTGACACCAAAATCCAAACATTTTAATTTTC GCATCACAGATTCATAAATAATCCACATTCTTTTCATGAATTATCCTCATTAGTACAAGC CACATGATTCAGAAGATTTGCAGTAAAATGCTTGGGCTGTGAAACTAAAGTCATTTACAA AACAGATTGGAATGGAAAATACCAAGTTCAGCTGAACTCACTTTAGCAGCCACAATAAAG TGAATTAACCCCAAATGCGTGATTACATAGAATTCTGCTTGAGCAACTCTCAATTTCCAA CTGTTAGTGTCTATAAACAAAGTTGTAAGGCATTATGCGTGCCATAGGCTACATCAAGTG AGCCATCAAATGAAGAGCTTGTCCTATTTGCTTAAAATTACAGAGATGCATGAAATCTGT TATGTACTTTTGAATTAGTAAGTGTAAGATTATTAGTGAGCAAATTGTGTGTCCTTGTCT GACTTTCTCAAGAAGTTTAAGCCTCATTAAAAGAATTAGCTAATGCATTGCTGTGAACTA TCTTTCATTTATCTGTCAGCATTTTTTCTAGTTCAGACCCTTCATATAATTCAACACTAA ATCTTAATCGTCATGTGCTTGTGTTAATTTATTTCACATTTATTAAGCACGTACTCTGTG TCAGCTATGGTGTGAGGTACTGAGGATGGACTGTAATAGATATTTGGGTCTGAAACTATA GTTCAGCTTCTCAGGGCCTTTGAAAGACCTTCTTGTTCCCAGCTCTTATCACAAAGTTTT ACAAATCAGTCTAGTGTGAGAAAATAGGCAGCAAACAAATTACAATTGCAGGGGCAGAAT CAGGAAGGCAGTAACTCGAGTCCATACAAAAAAAAAATAAGGAGCACCAGTAAAGGTAACT ACATAGGTAAATACTGTAGACAGAATAAACATATTTATCTTCTGTTATCTGATGTAAAGA ACAACTGCATAAAATAATAGCTATAAAATTGTGAAGATTCACCTTATAATGTATACAGAT GTAGTTCAAAAAAGGAGGAGGAAATGGAGCTGTATTGGAGCAAATTTGTTTTATACTATT GAAATTACATTGGCATAATCTAAGCAGCTTGTTTAGATTAAGTTGCTAATTTTAATTCCT GGTGTAACCACTAAGAAAATAATTTTTTGAAGAATGTAGAAATATAGGTAAAGTAACAAA AGAATTAAAATAGTATACAGAAAATATTTAACACAAAATAAAGCAGTAGTGAGGAAATAG ATTGACATAATAGATAAAAACCAACCAATAATTTAAAAAAACCCATGATCCAACTTTATG CTGTCTACAAGAGACATACCTTGTATTCAGATATACAAATAGGTTAAATGTAAAATAACA GAAGAAGTACTAAAATAATCACAAAAAGGGAGTTAATGTGGTTATACTAAAATTAGACAA AATAAATTTTAACAAAATATTACTATACATAGAGGGGACATTTCATAATGATGATGAG TTGATCCATCAGGAAGATATAAAAGTTGTAAACATACATGCATTTAGCCACTGAAACCCA

FIG. 1AI

AAATATACCAAGCAAAAAGTAGTAGAATTAAGGAGACAAGTAGGCAGCTAGACAATTATA CTCACCCCTGTAATCCCAGCACTTTGGGAGGCTGAGGCGGGTGGATCACGAGGTCAGAAG ATTGAGACCATCCTAGCTAACATGGTGAAACCTCGTCTCTACTAAAGATACAAAAAATTA GCCGGGTGAGGTGGGGGCACCTGTAGTCCTAGCTGCTCAGGAGGCTGAGGCAGGAGAAT GGCGTGAACCCGGGGGGTGGAGCCTGCAGTGAACAGAGATCGTGCCACTGCACTCCAGCC GCAGAATAGCAACAAGGAAATAAAAGATTTAAACAAACTATGAAACCACTGGGCTTAACA GATATTTTAGAACACTCCACCAAAAACAGAAGAATGCATATTTATCCCATTTGCACATAA AACATTTTCCAGGTTTTCTGACTAAAGTCAGAAACAAGACAAGTATGTCTGCTACAACCA TTTTCATTCAATGTTGAACAGAACTGATTCTTTTCAGGGCAAACAGGCAAGAGATAATAT TAACAATAATAAAAATAAAAGGCATGACGATCACAAAATAAGAGGTAAACTATTTCTAC TTGTAGGTTATGTGATATTTTATATAGAAAATCCTAACGAATTATTTTGCAAAAAAATAC AATCAATTTCATTTTTATACATTAGCAAATAAAAATTTAAAAATGAAATTAACAAAAAATA ATTTAAATAGCATCAAAATTAATCAAATACCTAGAAGTAGATTTAATAAAAGAAGCTTAA TAAGAGACTTCATCCAGGCTTGATTGCTTATGCCTGTAATCTCAACACTTTTTGGGAGACT GAGGCGGGAGGATCACATGAGGCCAGGAGATCAAGACCAGCATAGTCAACGTGGTGAAAC CATGTTTCTACTAAAAACACAAAAATGAGCCAGGCATGGTGGTGCAGTGCAAGACTATAA TCCCAGCTACTCAGGAAGCTGAGGCATGAGAATCATTTGAGCCTCAGAGGTGGAGGCTGC AGTGAGCTAAGACTGCACCACTGCACTCCAGCCTGTGTGGCAGAGTTAGACTCTTGTCAA AACAAAAAATTCTTCAGCATAAACATGTATATTTAGGGAATGTCCAGAAATTATAGAG ACATGGATTCCATGCAGCAGTTATAATTCCCTAAAAAGAGAATTATGAATTCACTGTATT GCTGAGGATTCTAACATAACCACCAAAGATCCAGGGAGAAAATTACCCTATTTTTGTATT TAAAAAGATGCATTTATTAAATGATGTGGTACTAGTCTCTATATAGGCAACAAAAATAAT GAAAAGGAAATAGCTCTGGATTATTAAAAATAAATAGTCTGTTAATCAAATCAATTAAAT AGATAATGTTCCTTCAACATTTTCAAGTCCTATACATGAATATCATTTACAATCATAATT ATTAGCAACTTCAATGAGTAGGCCACAGTTATACAAGTTTCTTGAGTCAGTTTGGAACTA TTTCCATTCAAGCAACATATAGTCCATTTCTGTAACATTTTGTTCTCCATCATTATATTC AGTCTCAGAAAGGTTACCAACACAGTCCTTGAATCACATGTAGTACAGGTTAAGCATCTC TAATCCCAAAACCTAAAATTCTAGCTGCTCTAAAATCCCAAACTTTTGAGAGCTAACATG ATGCCAGAAGTGGAAAAGTCCCCTGCTATCTCATGTGACAGGTCGTGTCAAAAGTCAACA AAAACTTTGTTTCATGCCCAAAATTATTAAAAATGTTATATAAATTTTGTTTAAAGACTAT TTGTATTGGGTGTTTATAAAATGTAAGTAAGTTTTGGGTTTAGACTTAAGTCACATCTAC AAGATATCTTTTTATGTATATGAAAATAATCCAAAAATCCAAAAAACTCACATCTGAAAC ACTTTTGGTCTCAAGAATTTCAGATAAGGGATATTCAATCGGTACACAACATATACACCT ACAATTACAAAATATCATTGAAAAAACTTAAAGAAGGACTACCTAAATTAAAAGATATTC TGTGTTTATGGATTGGAAGATTCAATCTTGTTAAAATAGAAATAATCTTCAAATTAATCC ATGAATTCAATACAATTCCTATGAAAATCCCAGATGGCTTTGTATTTTGGACACAAATTG TGAAAAGAAATGGAAAACTTACTTCCTAATGTCAAATCTTAACAAAAAGCCACAGTAAC TAAGACGGTGTGGTACTTCCATACAGTTAGTCATATAGATCAGTGGAATAGAATTCATGG TCCAGAAATAAACTCATATTTATGATTAATTGAGTATTGATAAAGGTTTTTAACACAGTTC AATGGCAAAAATCATAGTCCTGACAACAAATGGTGTTAAGACAATTGTATATCCACAAGC AAAAGGATGGAGTTGAACCTCACCTCACACCACATTCAAAACTTAACTCAAAATGAATCA TAGATTTATATGTAAGAGCTAATCTCTTAGAAGAAAACACAGAAGAAAATCATCATGACC TTGGCTTAACCAATAGGTTCTAAATATAACACCAAAACCAAAAGCAACAAATGACAATGT AGATACATTAGACATTATCAAAACAAAAACTTTTGTGCTTCAAACTGCACCATTAAAAAC GTTAAAAGTCAGCCCATATAAATGCAGAAAATATTTGCAAATCATATATGTGTTAAGGAA GGCAAAGACTTGAATAAACAATTCTGCAAAGAAGATATACAAATGGTCAATAAGCACATA AGAAGGTGCTTAACATCATTACTCATTAGAGAAATATTAATCAAAATCATGAGATACCTA TTCACACTCAACAGGATAGATTTGTTTTAAAAGGCTGTAATCATTATTGGTAAGGATGTG GAGTAATTGGAATCCTTCTACATTGTTGGTGGGAATGCAAAACGATGTAACTGCTTTGGA AAACAGTTTGGTAGTTCCTTAAAATCTTAGAGAATTACCACATTACCCACTAATTCAATC TCTAGTTATAGACCCAGAGAACTGAAGACATGTTTACACAAAAACTTACACATGAATGTT CATAACAGCATATAATTCATAGTAGCCAAAAAGTGGAAACAACCCAAATGTTATCAATGA

FIG. 1AJ

GTAAATGGAATAACTCATTGTTCTATATGCAAGCAATAAAATATTATTCAGCTACTAAAA GAAATGAAGCACTGATATATGCCACAAGATTGATGAATCTTGAAAACATACTAAGTGAAA GAAGCCAGGCACAGAAGGCCACATATTACATAATTCTATTTGCATGAAAATGTTGAGAAT AGGCAAATATATAGAGCCAAAATAATTTGTCCTTGGCACGGGCTGGCAGAATGGGACAAT GAGAAGTGACTGCTAATGGATTTGGAGCCTCATTTGGAGGTGATGAAAATGTTCTAGATT AGTTAGTGATGATGATAGTTGCACAACTCTGTGAATATTCTAAAAATCATTTTTTGAA CCCCTTAAAGCAGTGAGGTTTATGGTATGTGAATTATATCGCAATAAAATGTTTTCTTTT AAAAAGAAAGAACAAAAATGATGGGATATTTTAAAAATTTTAAAAATTGAAGACTTTTTTT TTTTTTAGAAAGTTCTGCTGCTGAAACCACAGGGAAGCAAAAAAGGTTGAACACAATT TGACATGTTAATGTAATGAGAGACTATAATAGGAATTATCCACGGGTTGTTTTATCTGTA CTTTCTGACTAAAGTTTTTTTCCGTACTTCTATAGACTTTAAAATGGTCCATAGATGTGC AAAAAATGAGAGAACCTATTCCATGAAACCATATATCAAGTCCCAGAGAGCAGAGGGAAA ACCTTTTTTTTTTTTTTTTTGCAAAGAAGAAGTCATAGACTGTGTGAAAGAATAATGT TGCGAGACAACAGATCTGGAGTTGGACAGGGGCAGGAGGCATAGTGAGAAGATCAGTTAT TGCAGTTGTCATCCATAAGGGCCATCTGTACACTCTGAAAGTGGAGCTATTCATAGTGAG AATGATGTTAAGAAAAGGAACAAATAAAATTACAGTCCTCGTTATAAGAATTTAGCATGC AAATCTTATCAGAGCAGTACTGAGGTAAACAAAAAGTGTCAAGAAATCATGGGATTTAAT CAAGGAAAATTTTCATGGTCCCTGCTGAACAGGGAAATGTAAGGGGATTATTGTTTCATA GAAGACCGCCAGTGCCTACCAAATATCTGTTATACTCTATTATGATGAAATGGGTAATAG GTTAAGGAATACCATAAGGGGAAAGGAGACTTGTCCTACAAGTTTCTTAGCACTTAGCAA ATGGAGCAGGCATTTGCTATGCATTAAAAAATAAGCATCATCCAAACTCTCAGACTCATC CAGCCACAAACTTAACTTTTTGTTCCTCCTCCTCCCAGATAAAATTCTCGACTTATTTCC ATTTGTCATCTTTTTCTCACTAACCGCCACCTCCACTGATGTCTCAGCCCACTTCAGTGT AGCTTCAGCTTTCATCATTACAGTGAAACAGCTTACATGAAAGTTACCAATGATTTCTAA AGAATATATTTTTAAAGTTTATTTATTGATCTTTTTGGCAGCATTAAGCAATGTTGTTT GTGGTTTCATTGCTCATATACTTTCTTCCTACTTTGATTTGAATACTTTTTTGCTTTGAAT ACTTACCTTCCCTGACCAGTAAATGCCACTTTGCTAGGTCTCTTCACAGCTCCAT GCTTTTTTCAGGTAGTCCCTTGCCCAGGTACTTTTTAAGTGAGGTGAGTATCAAATATA TATACACATCAGACTAGTCCTCTGGGATACACACAATCACAAATACACTTAAACACTCAA TGTACCTTTATTATAAATCTTGAAATGAGTTTTTATAAGTCTTGCAACCAAAGTTTAAAA AAGAATAAATTCTTTTTTTAAATTGCTTTTGGCTATTCCAGGTCTTTTTGCACTTTCATAAA AAATTAAAATTAGTACTTTCATTTCCAGAAAAAAGACTGTCGTGGTATTGAACGTGATTA ATTGCATTAACTCTATAGATCAATTTGGGGAGAATTGCCATATTAACAATACTAAGCCTT TTAATGCATGTCCACAATGAATATTTTTTTTTTTTGGAGTTCTTTTATTATCTCTCTGCAATG TATGAAATATATATATATATATATTTTATGAAATATATATATATATATATATATTT TATGAAATATATATATATATATATTTTATGAAATATATATATATATATATATATTT TATGAAATGTATGCCTAAAACACATTCTTTTGATATTGAAACTTTTAAAATTTAATTTTC CTCCTTGCTAAATTCTTTTATAAGTTCTAGTGGGTTTTTTGGTAGATTCTTTAGGATGATC TTTGTAAGCAATAATTTCTTCTCAATAGAACCAATCTGTAGGCATTTTATTTTTTTCT TTTCTTCTTGTATTGGCTCAAAGTCCAGTACAATGTTGAGTACGAGTGGTGAGAGAAGAC TTGATTTTTTGAGTGGTAAGCCAACACTGCATTGCTAGAATAAATCTGATTGAGCAAATG GTATTATCCTATTTATATTTGCAGGATTTAATTTGATAACATATTTTTAAAGAGATTTT TATCTCTATTCATGAAGGATATTTAGTTGTTAGCTGTCTTTTGTTGCCATATCTTTGATT ACAAAGATAAATGTGACCTCATGAAATTTGTTGGAACATATTATATTTTCTGTACATTTC TGGACCTATCTGGGCCTGGAGATTTTCTTGTAGCATAGTTTGTAAGTACAGAGTCAGTTT TGGTCATCTTTGTCTCAAGGGCTTTGTCCATTTCATGTAAGTTGGCAAATTCATTGTT TATCCATAATGTTTTAATGTTTGTAGCATGTTTGCCTCTTCCTCATAACTTTATCCTGG TCACAAACATTTTTTAAGACAGAGTAGGTTTTAAGGTCCATCATGTACATGCTATTTCCA ATTCATAACTGTGGTAATACATTTTTCAGGGTGTATTTTTGCATTAAATATGATTTATAA AGTTTATTCATAATAGTGAAATAAAAGTGGGGTGCATGTATTTTACTTAATCCTTCTCAG TGCCTGCTTGATTGAAACCTCTGAGATTTACAATAATGTACTTTTAGGGATGCATTAAGG ATTACTAGTGCATAGTTCCTGGAGCTCAGTAATGTCAGTTATTCCTCTTAATTTTATACG GAGTTTCTCTGAATTCTCCATGTCTCTAGACAGCTTATCAATGGAGAAATTTATGTGTCC

FIG. 1AK

TCAAAATGAATGCAGGATTCAGCATCTTCTATCCTTATTTAGATCATTATCTAAAAAGGG CATCACTACATTTTTTTTCCCGATTTCAGGGACCATAGCTTTCTCTTTTATGAAAACTGT ATTTTTTTTTTTTTTTGAGATGGAGTTTTGCTCTTGTTGCCCAGGCTGGAGTGTAATT GTGTGATCTCATCGGCTCATTGCAACATCCACCTCCTGGGTTCAAGCGATTCTCCTGCCT CAGCCTCCTGAGTAGCTGGGATTACAGGCATGTGCCACCACGCCCAGCTAATTTTGTATT TTTAGTAGAGGCGGGGTTTCTTCATGTTGGTCAGGCTGGTCATGAACTCCCAACCTCAGG TGACCGAAAACTGTTTCTAATGGCGGCAGAAGTCATCAGATGCAGAATGTAGATTCTCTC CTTCAGGGGAACAGTCAGTGATAGAATCACTAAAATTTAATTGATCTATCAGAGATCATT TAGAAGACAGACAGTTCAAGATCATTTAGCAGACACATACAGGCTTTTCATGATAGGAGT CTCCTGGAACATTCCAGCATCCATTGCTCATTCTTTTCAGTTATTTTTTTAAAATTGCTTT TTAAAATGAGAGTCACAGAAGAGAAAGTTCCTATCTCTCCCCAACCAGTGGGTTAAAAGA TTGAGTTGAACCACTACTATGTAAAAAAGATTGTCTACATGACAAGACATACAGAGTGAG AAGAAAAATAATTTATCCGATATTTTCCATTCAAGGGCAGGTCTTTGTTAACATCATTTG CCTCTTCAAGAAAAATGGTCAAAGGAAATGTCATATTAATTTATCTGTGTGGACATA TAAGTAAAATTCTGTTCTCAAATTAAAGATTATCGAACAGACTTTGATCTGGTGGTGTAA AAATCAACAAAATCTATCGAACATCTATTCTGAGAAACCACAAGGACACATTGGTCAGTA CTGGTTTGCCGCACAGAGACAGAAAGTAAAAGCTGAATAATCTTAACAGAGCTAAGGTGG TTGTTCTGTTCACTAAACCCTATCAATCTTCATGAGCTTACAATTAAGAGAATATTGTAC TTGGAGGGATTCCCTGCTATTATCAAATAACTTTGAAAAGAAATGGAAAAGTACAAGTTG TGTAATTACTGTTACAAATTCCAGCTATTTGAAATATTAATGTAAGACCGCAAAAAATCC TCAATGGGTTTGTGTGCATTTTAAAGGGCTGGACCACAAAACTGATTTCAAACAATTTCA AGGTTTGCTTTGTCCGAACTCGAGAAGCAGAAAACCTGAAAAGCTACAGGTAGTTAAGTT CTATCTCTCTGGCAGCAGATGGCAGTATTGATGCGTGAAAAATCCATAACAGGTTGCTTG ACGTTACTTGCTGGGTTTTCCTCTGCTTTAAACTTTGGTATCTGAGCTGAACAAAAATTC CTAATAAGATAATATGGCTGACATCCCTTTATCATTCTCCCTTTCCCAAGCTTTGTTCTTT TTACAAGGAAATATCTTTTCCACTTGCAGCTTTCTTTAGACATTGACAAAATTTTGATGT TTTAACTTTTTTTCCACACAAACTCCTATTTGGTATTCGTCTGAATTAACGCCAAGCAC ATACTAAGGTCAGCAAATGCTCTGGAGAAACAGGCGCTCAAACCTCCCACACCTCAGGCG ${\tt TCTGGAAGCCTTTCCTTACTGTGTTTTCTAATTACTTCCCCAAAGTGGAACTTTCCTAAG}$ TCAAATTGCAATAAGGGTCTGTCTCTTCTCCTTTCAGATCCCTGGAACATCATCTGTAGT TCAGAGAAAATGGAAGCCCCTGCAGCCTGTTTCACAGCCTCGAGGGCCAGGACAGCCAAC GAAGTCCCGGATGAGCGCTGTGGCGGCTGAAATAAAGCAGATCCGAGCCAGAAGGAAAAC AGCCCGGATGTTGATGGTTGTGCTTTTGGTATTTGCAATTTGCTATCTACCAATTAGCAT TCTTGATTCTTAATTAACTTTTTTTTTTTTTTTTTTAACTAAGCCAGAGAAAAATCTAAACT TTCTGCTTAGATACCTTGTCAGGCCAGATGACTCAGTTATGTTGTTACCAGCAGGTAAGG CGAACAGCCTTTAAGAGTGCTCAGACATGTGCTTTTGTCATGCGTATTCTCAGTTGCATG GCAGACATAAAACAGATGTTTCTCCAATCTCTTCAAGCTAGTTGCTAAACCTTAGATGCA AACGGGTTTTTCAAAACTTCGTTTCCAAAAACATAGGCAATTGTGAGAGAATTATATCTT AAGGATAAAAAGAGATAAGAACCTTATGTTAGTATTCTAATTATACTTAAAAGTGCATTG GAATTCATTTCCTGAAGTCACTGAATGAAGCTCAGGGCAGATAGTAATAAAAATCAATGA GGGAAAGTATGCTATTTGCTACAATGCAGGCACAACTATTAAGTTAAAAATTTTGACCCA TATTGGGAATTATCGAAACTGTTCCACATAGACTGGTCCCAAGGCAATACCAATTCTTGT TTACAACAGGCTTCGAACTTAAGCTAGAATTGCTCCTCTCACTTTGGCCTGATTCAGAAT CAATATTATATTCCTCACAGCTGGGAACTCTGAAGAACAGCAGCTTTGGCTGGAGTCAGA AGAAGTGGTATAATCAGCCGCAAAGGGTTCTCATTCTCTTTGGCCCCTGTTTTTGATGGT TTAACGGCTTTTTCAATGGAGAAAATGGAGAACAAACTTCTGTTCAGTGATACTATTT TGCCAGCTCACAAGAAATAGAAATTATTTTGCCCTCCATATCTTTTTGTGCTTTC TATTATGGCCACCTGCTAGATACTCTCTTCTGCTTTTAAGAACTTTATAAACATGTTATG

FIG. 1AL

39/51

TAAATTCAATGAAAATTGGATTTACCTCTCTCTGGGTTACGTATTTGGATTTCCTGTATC TTCTTCTGACTCAAATCCCTTCCTTTCCAAGGAAAACAAATAACTTTCAAAGGAGCAAGG CTGTGTTAAATTTAAGATATTTCAAGTTTTTGGGGCATTCTACTCTTTTCCACAACATAAA CATTCCTAGCTTTTTCTCCTTAAAACTTAACTTTTTGCCAAATTAGTCAAAAGCAATTTC TTTACAACAGTTCAGGTTTGTCCAAGATTTCAAAGACATTTTGAGGTAAAGGGTCATAAC ATAGTACAAATTTCTTTTGTCCGTATTATTTCACTCTATATAGTATTTTTTGTAAAACTCT AGTACTCTTTATACCAGAAATGGTATAAGGTACACCTTATACCAGAAATGCATTGTTGTC ATCTTATTGAGTAGCTGTAAGTTATCTGTTATAGGTTAGTGATTAGAATTTATAGATGGA ATATTTCTAAGTATGGAGAAAATTTTTTAATAGTCTTTAAGGATAGCATAACAAAACATT TTTTAAAGTTTAAAATAATACATGAAAAATTAACACTCATTAATTTTTAAAAATTACCAA AATTCTGCCCATCGAGAACTGTTTCTTCTCTGGGTATTAAGGAGTCCCAGAAGGCAAGTT TCAGATAGTCCAGGAAGATTGGAGTTGAAGGCATATGATACTTTGATCAATACATAAATG AAAGTAGGAAGAAGTACTTGAAGACTATCATTTAGGAGTGATTTTTAAATGATACACATA ATATATTCAACTTCCATATTCTTATTTACAAGAAATACTGCATCTGCTATTTGTGGGAGG GTTCAGGCACTGAAGTTATTCTAAACCAATTATGGGGTCAGAAACCAATCTGTGGTCAAT TCCTGCAACTGAAGAGGACAGGAGTCAGACCATCCTCTACCAATAGCCTTGTTCACCTTT GAATTTAATTATTTAAAAGACACTTTTCTGTTGTTTCTTTTCCTGCAGAGTATTTGGGAT GTTTGCCCATACTGAAGACAGAGAGACTGTGTATGCCTGGTTTACCTTTTCACACTGGCT TGTATATGCCAATAGTGCTGCGAATCCAATTATTTATAATTTTCTCAGTGGTGAGTTTTC AACTGTTCTTCCATAAGCCACAATTGTAACCAAGGATGAGGAATCAATGAACACTCTTCA ACTATATGAGGAGTTTAGTTGCTATGTGAGTTGTATTTTTTCCCTGACCTGATTTATCTT GAGTTTCTTCTCTTTTGAGGCAAAGTATTTGTTACTGAACTCATCAGAGAAAATGAACTG ATTTTTCCATGTCAAACGTATAAGAAATGTTATAATAGAAGAAAAGTAAACATTCTGAGA AATCAATAACACAAAATCTTACATGACATACTTTAAACTCATGATTTACAAAAATATAAA ATACTTTGTTCTGTTTTGCCTTGCTATATTATTCCTTTGCCAAAATGTGTAGCCTAATTG AGACAGAATTGGGATCTATTCACTTTTAGATATTTTACTATATTTACGTTTCTCTTGTGA GTATCATCTCTTGGATTTATCTCAATATTTCCCACTGACTACCAAAAATAGTATTACTCC AAAATAACACATAAGTTAAATGATACACACATACATATACGTGTAACTTATACAATTTGT ATCTGTTTATGGAATCAATATAATTATAAAAGTCATTTAAATCACTATTGTTTATTCACA TTTTGCCCGACTGACTTTTAGAATTATTTTTAATTAGCTACCTTTTTACATTGCCTTAAT CTCCAACTCATTGGCGATTTCTTTGTTATTTCTATCTTCAAATATATGGTGATTTTATGT GGAAGAATAGAAATTCATTTTGTGGCATATTTAATAAAGCTTCTGCATCTTCCAACTTGA TCTTTGGCCTTCTGGTTTGCATAGGTTTAAAAAAAAAGGCAACAAATTAGATTGATGAGAA ATAATTTTGTTCTATTTAAAAAAAATCTAGCACAATGACTAAAGCTCTGAACCTCGCAC TAAGCAGGTAAAGGCTATGAGGAAGTTGTAATGAGAAGTGTTTGAAGCAGAAGTCACAGA ACCAGGTCAAAGTCCTAGTATGGAGGATAAAAGTGAGTTAGAGGAGGCAACTGATAATCA CTGATAACTCATTATGTGACTGCTATTGTGCTGGGCCCGTGAACATTCATCTTCTCATTT AATCACTGATAACTCAGTGCGTAGCTGAGGCTAAGAGAGAAAATGATTCGCAATACTG CCATTCACACTAATAAAAGTGATTCATTCATTCTCATAGTTCTCCAATATCTCCTCCATA ATATAGATTATGAAACTGAAAATTTTCTGGATATTTTGAGTGTATGTTTCTAGGTATTTTG TGGATTTAATTGTTTCAGTATCAGTTATTTAGAGTAAAATGCAGGAGTAATTTTTGTATA ATTTTGGCTTTGTATGACATAAGTTTCATTGTGTTTAATTATTAAATATCTCTGAGAGTT CTTCTACTGATGATCACTTCCATTATAGTTATGTAGATAAAATATACCAATATGCGTAAA TATATGAGGTTTGACTATAAAGGAATGAAGCCAAATTCCAAGCCCCATATGTGAAAGGCAG CCTCGTTATTTTATGAAAATATTCATTGTTTCAAGAGTCTACCAAGCTTCCAATAAACTC AATTTCCTTATTCTATTTTACCCATCTTTGCAAAATATTACACCTCATTGTTAGTTTGGC TCAAGGGAGCAACTCAGTTGTACCCTATTCATAATTTGTTGAAGCATTTATGTATAATTC CTTTTCCTTTCATTCTCTGTTTGCCAGGAAAATTTCGAGAGGAATTTAAAGCTGCGTT TTCTTGCTGTTGCCTTGGAGTTCACCATCGCCAGGAGGATCGGCTCACCAGGGGACGAAC

FIG. 1AM

TAGCACAGAGAGCCGGAAGTCCTTGACCACTCAAATCAGCAACTTTGATAACATATCAAA ACTTTCTGAGCAAGTTGTGCTCACTAGCATAAGCACACTCCCAGCAGCCAATGGAGCAGG ACCACTTCAAAACTGGTAGAATATTTATTCATATGACAAGGATACCTGAGTAAAACTATC CTTTTTAAAATCACTGGGAACAGAAATTTTATTATCCTATGATGTGAAGCTAAAATTACT **AATGATTTCTCAACTTTTGATTTAAATATGTTAGAAGTTTAACCTTCAATTGAGCTTATT** TCAGGCTATTTCACTTTTAGTTTCATGTATTAAAATGTGTGTCAATTAAATGTTTAAACA TTTCTAATTCTTTTTATAATCCCTTGTTATTTTAATCTCTCACATTCAGATTGGTTCCTA AAAATTACCAGAATCTATCCAATGATTTTTTTTTGCTACTAAAAGAAGTAGCAATTACTAA TTTGCAAATTGACCATTGCTAACATTTCTTGTCTCAACATATGGCCAGTAAGACTCTATC ACAGTAAAAGTTTTAACGTAATTTCCATCTCTAACACTTTAACATTTAAGAATAAGCTAA ATCACATCATTATATTCTTTTAACAACAACAACAAAAAGTGATATAGTCAGCCTTGCTGG ATTAAATTAAAAATGCACCACTGTGCTAGGTGCTAGGGAATGAGATGGCGTCGATGCAAA CATGCCTTCAAAAGAGCTTCAGTCTAGTGAGGGAGACATGTTGACAGAGTGCAAGGCAGC AAACAATCTGGGGGACAATTCTTGGTCATGGCAGAGCAGTGAAGCTACCAAGGACAGTGG TGACAGTTGATGTCAGTGGATTCCATCTGGGCCTGGGGCTGAGTACCAGGTGGTTAAAAA ATAGAGGGGCTTGCTCTTAACTCACACATACATGAATAGACTATCGTATATTTTGTAGAA AATGTAAGATCTGGGAGTCAAAGCACTGAGTATTCAAACTTATTCCCCTGAAAAATTCTT CTGATTCAAATATTTACTTGAAAATTAAACTAAAAGTAAAAGAAGTGTTTATGAAAGATG ATTTTCATCTCCTATTATGGTAACAGGTGTTCTGATTGTATTGAAACAAAAGATATGGGG CACAGTGTTTAAGAAAACTTTCATAGAAAATTAATTTTTGTTATTTTTTCATTTTTCCA TTACACTCAGAGAAAAGTAAAAGAGCCTAATTATCCACAACTTGTTTTCAAATCTTGGAA TTTGGGATTCTGTTACCTTGTGCCTTTTATGACTCAAAGCAAAAACTATCTTCTTATACA AGGTTTATTGAGATCATATTGTAAAATATCAGCACTATATCAGTGAAAGCAAGGTATTTT AAAGGAGAGTTGTAATTTTCGGATCGTGATGACAGCACTTTAAAAAGTTTGAGGATAACT TCAAATAACGTTGATAATATGCCTTAATAGCCAGTAATAGCTCAGAGGAAGAGTAAATTC GGAAAGGGGGAGAAAAAAAACTGACTCAGAGAGCAGCAGTTATGTGACGTATGGGA AGTCAGAATTCCTTTGCTCTAAATCAGTGATTCTCAAAATGTGGTTTCTATACCATCGGC ATCAGCATCATCTGGGAACTTAGTGGACATGCTAACTTCCACCCTATCCCTCACCTACTT AACCAGAAACTTTAGGGGTGATAGCCCAAAAGCTGTGTGTTAAGCACTACAGGTGTTTCT GAAGCACTTTAAGATTTGAGATCCACTGCTTTAAGTGATACCATCTGACATCAGTTTATC TGCCTGTGTGAAATAAAGTCTTTTACTGCACAGGTGTCTACAACAGGGGCCACCATCATC GCTACCGTCAACGTGGTTGGATGTCTGAAAGAAGAAGCTGAGTATCAATGTTGACTCTCA CTCATGTCATCTTATTAAAAAAAAAAACAGTTTACAAAACAATTGCTACTGATAAATGCAG TGTGAAAGACTGGTTTTAAGGCACTTGTGTGTGTTTATGTCCACCCAGATAACTTGAGTTT CAGATAGTTTCTAAGAATGATCATTTCATGGAAGGAGATATAAAATAAAATAAAACCAGT ACTTAAACTCTGGGAATGTAATAGGCCATGTACATAGCACTCAACATGTGAATCCAGGAA TCCTTCTAAGAGGTCTAGATTTAGTATGGTTACCTTAATAGGACAAATGGTAAAGAAATA GGTGTTCCCAAACTCTGCCAATCTTATGAAACAAAGAGTCAACTCTTTACCTCATTATTT GCTAATGACACAAATGCAAAGACATCTTTTGAAAAGAATGTGTTGGGACTGTTTTATGCT GTACCTTGAATGTGTATCTCTCCTTTTTTTTGCTATATTTCAAAGATTTAATGTAAGTTGT CAATGTCATTGAGTTCTTGTTATCAATAGGGATGATATAATTTTATCTAACATGGAATCC ATTTTAACTTTGTTATTTCTGAATTTCTATGAAACCACAAAAACCTTCATACTTGAATTT AGCTTGGAGAATTTATTGGATATTAAATACCTGTTATAAATTATTGATGAGTTAATTGCA AGTAGCAGACACAATGATATTGAATTTCACTCCCAATACACATTGTTTTAATGAAGATTA AGGTAAATATGTTTATAAAATTTAGTCTGGCTATGCTTAAACCTGAAATAGCAGAATGGC AAAAAACCCCAAAGCTGTTTATGGACCCAAATTGTGAGGAGGGCTATTATTTTAATACTT GTGTAATAATAGAATGCACTTGATGTAAATTGTAATAGCCATCAACTGCATTTCAAAAAC

FIG. 1AN

CCTTTTCTACTTCTACAAGTCACAAGTCAAAAAAAAGTAAATTCCACCAAGTTTTATTC AATTAGTTTTCAAATTGCATGAAGCAAAAAATAGATTTTTAGAGACAATATATAAATAGA GCATGAAAAACACTATTATTTCCCAAAGTTCAAAGGGAATTGTTTTCTACGCAACTACTG CTACTAACAAGGGGACAACAACCCCCTCCACTTGCCACGTATTTTTATTCTCTTTT ATATCTTTGGAGTTAAATGTCTTTTATGTTTTTCATGAAATGTATTCTATAATTGTTGTA TTTCATGTGTGTAACATTATGTCAGTTGTTTTAACAATTATCTTATATCTTGAAATTCTT TATGCCTGATTGTACTGTGTCTTCATGAAGAAATTTCTTATCAAATCCAATGTGATTACA CACTTACTGCTGTAAAGGATGCGCATTATGTAGTTTTTTAAGTAAAAACTATAGTGAGAAT TCTATAATCACATTCACACTCCCCTCTCTATTGTATGAAAAATCTTGTTGTTGTTGATTA GATAAGGTGGATATTCACTCATAGTTAATGTCAAATCTCTGCAGTTAAGGATTGAATTAA GCCCTCTGGTGCAGTACCTAATGATCAAAACATTTTTTCCAATAAGTTTATATAACCAAG GATAATAATGATATAAAAGGTTTTTTAATGTTGTTTTTTAAGAGCAGGTACTATAACAAAGA AGGTTAACACTGGTACAGAAATATTTCATAAAAGTTATGAAAACCAGATAAATACAGTAT TAAATTTTGGAGCTTTTATCTGAGTTGAGAGATTTAGTCTACATTGACTGAGATGAAATG TTTTCTGGGCCATTTTGTATAAGTCATTTAGGACTATTTTAAGTTCACTGGTAAATTTTA AAATGTATATTTTCAGCTTTTCAATTTTTTTCAAAATAGTTCTGAGAAATTACAGAATCA ATAAAAATGAAGCCAAAGTAACCCGTCAAGGTAAATACTTGACTCCTAGGAAAATGTGA TTTTAGTAGGCATCTCAAGAGGAAGTGAAACTTCTCGTGGTGAAATTACAAGAAAAACAA GTTATTCAGTGGTGAGAATGTGTTGCTCTAAGCAATCCATTAGCACAGACTAGCTACTTG GCCACTCCTCTTCCTTCTGGAGCCAGCCCTGAAGAGTGGTCACAGCATCTTCATTTTAT CCAGGCCAATGGCCATGCATGAGAAGTTGGGTAGCAAAATTCTTGAAGCACCTCTTTGTT CTTGCTCTTCTTCACTGTTTTCTCACTCTCCACCTGTAATGCTCACTGCCAGTTTTACC ACCAAGCTAAGTATCAGCAGACCTCCCTCCACAGCGTGCCTTGCCCTGTAGAACTCCTGG TCCTTCCTTCAGCCCAACCCCATCCAATTGCCTAGGTTCTTGTTGTCTCCTGAGATGAAC AAGAGGCAAGTAGCTAATTTGAGAACAAATGAAGCAGAGCTGAAGGAAAAAGTAAAACAT TCCCAGAACATTTATGCTTATCAGTGGTCTTCTGAATCTGTGACAACTCCCTTTTCAAGC CCCAGCTAAGCTTCTTGCCTCAAGCCAGAAGGAATCCCAGTTTTGAGTCTTGTGTTAAGG CCATGGCAGGTCAGTAGGGAGATTATCTGAGGAGGTACCGCTTGTGACACCTTCAGAAAC AAAACAGCTATTGCCTTACGTTTCATAGGCCCAGGCCCTGAGCAATAGCAAAAAGATAAT AATCCTCAGTGCTTGACAACATGAAACTTATTTAACTTATTATAGATGAGATAATGAGAA CATCTTCAGAAAAGAAGCTATGTTCCTTAAAACAGGGGTACAGATTTAAAAGCTCTGTTT ATATGGTTTTGGTAGACTAAGTGAAGAACTTGCCTATAAAGCTGAGTCTCGATCATATAG CATATCCATTATAAAGTGAGAAAATTGCAATTTTAGAGTATTGTCAATACATCCAAAAAT TTTTACATGATTTCTAAATGCAGATGTGTGTGTGTGTGTTGTCTACGTATGTCTCCCATA TGCAACAAGCAGTTAATTAGTCCAAATATATCCCACAGTGTAGATTAGTTTCATATCTCA GCTCTTCAATGTCTCTTCTTCATTTAATTCACTCCTTGGTGTCTAGTTTTCCTCACTCTT TTACAAATATCCAGGTTCTATATTTCTGCTTTTCTAGAGAGCTTTTTCCCTCAAGAATAT TATAACAATTTGAGACAAGTGAAAAGGAAAGATCTGTAAACTGCCTATCTCCTTTGAAAT TCATTGCCAATAATCCTTAAGAATATAAAGTTCCTTGATGCCAAAGACCTTCTCATTAGT GTTGCTGCCTGTTGTTTCATTGGTTCCCTAGAACAATGCCTGGCACATAAAAGTTATTTG ATAAATATCTCTGCTATTAATGAATTAATAACTGCATGACAATTCTTTCCTCAATTC ATCATTTTGCTTCATTTTCTCACAGTTGCTTCAATGTGTCTGTGGAACTATCTTTCCATG TGAACAAAACACTCTACATTCTCAGTGTCTACAAAGCACATATTTCCTTTTATTAAAATT AAACTTTGAGAGCACCAAATCCTAATGTCTAACCATCATCAAACTGGCAGATAGCACCAG CTGACACCTGTAAGAGGATTTATCATGGTAAACTTCTCTTTGTTACTGACATTTTCAGCC TCTTGGGCTCTCCCTCCTTACTTATACACATTGGCACCCAGCTTGAAGTCATACTCTCT AGACCCTGGGTCAATGTGGGTAATGCATCCAGGAATCCAGCTTAACTCTTCCTTGGTCTC TTTGATGTGACTGACCTTTATTTCTACATTTCTTCATCAAACCAGTCTCACAGTTTTGCA CAGTGCAAATCACATGCTGCACCATGTGCTTATTATCTCCTATAACAACAGATGCTCCAC

FIG. 1AO

TGAAATGCAAAACTCTGTGTTAAGCCAACAACTGCTTCTCCATCCTTTCCTCCTATACGT TTCTTCTCACTACAACTTCCCTTCTCAACCCCAAAGGGACTACTGGATTCTTTACTCTTT TACTATTTTGTCCTGATGAAATTCATGACAGTTTTCATACAACAGAAAGCCTGCCCTCTT AGAAGAGAAGAACTGAAAAGAAATGGTTGAAGTAAGGTAGAAAGCCCTCATGGAGTTA GGTGGCTAGGCCAGCAGAGCTAGGCACTGTTCTCCTGTTCAGAATTGCACTCCTGATACT CCAGATGGGAAGCCTGCCATGGCACTAACCACAGCACTTTTTATACCCTATCTCTGCTAT TATGAGCCCACATTAGTTTTTCTTCTGCTTCAGAAATTGTTGCAAAAAATAATTTTATTA TTTACAAATTATTTTTAAACCATATAAATCTGCTTAGTTTGATTTCTCAAACCCTCTAAA ACTTACACTTCTTGTTGTCCAATCTTTGCTTTTAATTGGGTATAATTTGAGGCAGAAATA **AATTAATCTCATTTTTAAAAATGTACTAGCTATTAATAATTTTTTAAATTTTATCTTCTAAA** CTTTATTACTTCCATTGAGAACTGTTAAAATAACAGAACTTACCTCACTGTACGCTGGCT TTTGAAAAGGCAGCAGAACTGTTTATCTGATTATCGAAGTAATCATATTACATTTCTTTT TCTTTTCTAAGAGAAACCTTCTTCATGTGCTCAGTCAAACATTTTGGTGTTTAAGAATTG ACTTATTAGGTCAGGCGGGTTGCTCACGCCTGTAATCCCAACACTTTGGAAGGCCGAGG CAGGTGGATCACTTAAGGTCAGGAGTTCGAGACCAGCCTGGTCAACATGGTGAAACCCCA CCCCTACTAAAAATGCAAAAAAAAAAAAAATAGCAAGGTGTGGTGGTGCACATCTGTAATC CCAGCTACTTGAGAGGCTGAGGTGGGAGAATCATTTGAACTCGGGAGGCGGAGGTTGCAG AAAACACAAAAAACAAAAAACAAACAAAAAAGAGTTGACTTAGTTAATGAAAATATTTTT ATTAGGAAATTATACTTCTCTTTACAAAGTATGTATTATTTGTTGCATCTATATAGTCTA TCAATTCTAAAAGCACACTTTATGCGAAAATGTAGTCTAGGCCTTCAGAATGTATTATTA CAAGAAAGTATCTATCAACCATGTTTCATTTGTTTGCATGTTTTGTTTTGCAATAG **ACTATGAATATTCAGCTTCAAATGCTACCTCATGATTGTTACATTCCTGTTGTTGAAAGA** TCATTCCATTGGCAGTAATCTGTGATTCAAAAGTTAACAACATACCATGTATTCTTGTAG GAGATTATTTCATGCTTATCACTGATCAACTTACATGCAGGTTAAAACCAGCCCTGAAAA AATGCTCATCATCACTGGCCATCAGAGAAATACAAATCAAAACCACAATGAGATACCATC TCACACCAGAAGAATGGCGATCATTAAAAAGTCAGGAAATAACACTTGCTGGAGAGGATG TGGAGAAATATGAACACTTTTACACTGTTGCTGGGAGCGTAAACTAGTTCAACCATTGTG GAAGACAGTGTGGCAATTACTCAAGGATCTAGACTAGAAATACCATTTGACCCAGCCATC CCATTACAGGGTATATACCCAAAAGATTATAAATCATGCTACTATAAAGGCACATGCACA TCAATGATAGACTGGATTAAGAAAATGTGGCACATATACACCATGGAATACTATGCAGCC GGAATTGAACAATGAGAACACTTGGACACAGGGTGGGAAACATCACACCCGGGGCCTAT CATGGGGTGGGGGTAGGAGGGAGGGATAGCATTAGGAGAAATACCTAATGTAAATGATGA GTTAATGGGTGCAGTACTCCAACATGGCACATGTATACATATGTAACAAACCTGCACGTT GTTCAGATAACTGGAGCCATCTTCCTAGCTCTTTATTTCTCAGACAGTGTGGGTAAGTCC TGCTCCGTACGAATGCTTATGTCAGTTTTGAAGTTCAGTACTTTCTTAAGAGCCAGAGTC AGTCAAGATGTTCCCTTAACAAGATTTTTCAATGGGGTTACACATTAATGAGTTCTTTTT CCTCCTTTAAGTATTTGAAAATTTTGGTTTAATAAAAGGTTTAACTATGATGAATTTAGG TTCATGTTGCCACTAGGAGTGTCCAGAATAGACAATTGAAACAGCCTTCTAGCTACTACT **ATCAAAAAGAGCTTTAAATAACATATTTTAATTAAATAACATTATTTTCTATAGCTATA** CCTCAATAAAACCATCAACCAATGTTTGTACAATTTGATGCCCCCACTCTAAGATTTTTA GCTAGTGTAAATCAGAGTCTCCTATTTAATGAGACACTTTATCCAATCAGGTTGTGTTTA TTATTCAACCAGATGATCTTGGAACTTATAACAAACTAGTAATACTTAAAGCTGGGCTTT ATGTGCGTGATTTACTGGGATGTTTGCTTATACCTTGTTTCCAAGCTAAAAATATTGTGA CCAGGTGTGTTAGTCTGTTTTGAGTTGCTATATAGGACTACCTAATGCTGGGTCATTTAT AAAGAAAAGAGTTTATTTGGATTATGGTTCTGCAGGCTGTACAAAGAGCATGACATCAG CATCTGCTTCTGGTAATGCCCTCAGGAAGCTTTTACTCATGCCAGAAGGCAAGGGGAGCC

FIG. 1AP

GGCACCAAGCCATTCATGAAGGATTTGCCCTCATGACCAAATACCTCCCACTAGGCCCAA CCTCCAACACTGGGGGTCTCATTTCAACATGAGATTTGGAAGGGACGACTATCCAAACTA TATCATCAGGATTTTCTGGCATGGACTACCAAGCCATTTCTGCTTCAAACTCCCCTGAAA TTCTTGTTAAAAATGCAGATTCTTTGATACCACCCCCAATACACTATTTAGTCTGAGATG AAACTCAAGGATTCTGATTTAATTGATCTAGACTAGCATTTGACCATTGATTTATCATCT GGGATTCTAGGAAGTCAACCACTTATATGTTTTAGAGCAGACTTCATTATAATTGAGGAG AATGTTTGTAGTCTGTGGGCTCCTCTGTCCACTTCTGATTGGGGCCCCTTTGCCTGATTC TGACTGGATCAGGCAGAGTTTTATTCAAGCCACTGTCCTTTTTTGGCTTCTTAATGTTCAA AATATATAACACAATCTCAGTTTTCTAAGAGCTAAATTATACGACTTGGTTCTTGTCTG GTAACATAACTGCATTACTGGATCTTGTCAAGATTCAGAGACATTCTCCCAGTTTCAAAT TTGTAACTAACACTGTTTGATCACAAAAAAGTTCTAAGCCAAAGCAAAACTCTTTCTACC ACCACCAGATGGCGTTACTTTGGACTTACCTATAAATGGATTTCCAAATGGTTTTTCAGA AACCAACTGGAGGTACTTAGAAAAACTTATGGAACTCACAACTATTCTTTGCATGTCAAA TCCTCCAGCTTATTTAAACTATATCCTTATATTAACCCTTGTTGGAGATGTGTCCTCTTA TTGCACTGTATGTGAGTGTGTGTGTGTATCCCATCACGTTGGTATGATGATAGCACCC TTCATTGAGAAGCTTTGCAAAAGAATATAAGAACATGTTATTATGTTTACTTAAAAGTA TAAGGCCGGGTGTGGTGGCTCACACCTGTAATCCCAGCACTTTGGGAGGCCAAGGTGGGA GGATGACGAGGTCAGGAGTTAGAGACCAGCCTGACCAACACGGTAAAACCCTGTCTCTAA TAAAAAATACAAAAATTAGCCAGGTATGATGGCACGCATCTGTAATCCTAGCTACTCAGG AGGCTGAGGCGGGAGAGTCCCTTGAACCCAGGAGACGGAGTTTGCAGTGAGCCTAGGTGG AAAAAAAGTATTGAGGACATTGCTCATGACATTCCAAGGTTATATAAAAGAATATATAA AAAGAAATTTCTGCCTGGACTTAGTGCCAGGAATACTTGTACTTTTCTTGCTTTCTTCTT GAAGGATTAGAGGCAGATCTGTAGCATGCCGAGTCCCATCTTTGCATACAGGCTATCATG ACAAACATTGTATGTGCTAATTCTATCTGGCTTCTCTTTATATTCCTATCTGTCTCTATT TCCTGTCATTTTAATGTTTTAAAATTGTACTTTTACTTAAATGGTTTTTTGGAAGAAATA AATATAAGTAAAGTCTGTTAGAGGCCCGGCGCGCGCTGCTCACGCCTGTAATCCCAGCACT TTGGGAGGCCAAGGCGGGTGGATCACAAGGTCAGGAGATTGAGACCACCCTGGCTAACAC CGGGTGCCTGTAGTCCCAGCTACTCGAGAGGCTGAGGTGGGAGAATGGCATGAACCCAGG AGGTGGAGCTCGCAGTGAGCCGAGATCTCACCACTGCACTCCAGCCTGGGCGACAGAGCG AAAAGAACGGCAAAGCCAACAAACATATGAAAAAAAGCTCATCATCACTGGTCATTAGAG AAATGCAAATCAAAACCACAATGAGCCATCATCTCACGCCAGTTGGAATGGTGATCATTA AAAAGTCAGAAAACAACAGATGCTGGAGAGGATGTGGAGAAATAGGAACGCTTTTTACAC TGTTGGTGGAGGTGTCAATTAGTTCAACCATTGTGGAAAGCAGTGTGGCGATTCCTCAAG GATCTAGAACCAGAAATACCATTTGACCCAGCAGTCCCATTACTGGGTACATACCCAAAG ACAATAGCAAAGACTTGGAACCAATCCAAATGCCCATCAGTGATAGACTGGATAAAGAAA ATGTGGCACATATAATATACAGCATAGAACACTATGCAGCCATAAACAAAGGATGAATTC ATGTCCTTGGCAGGGACATGGATGAAGCTGGAAACCATCATTCTCAGTAAACTAACACAG GAACAGAAAACCAAACACCACATGTTCTCACTCATAAGTGGCAGTTGAACAATGAGAACA CATGGACACAGGGAGGGAACATTACACATCGGGGCCTATTGGGGAATGGGGGCTAGGGG AGGGATAGCATTAGGAGAAATACTTAATGTAGATGACGGGTTGATGGGTGCAGCAAACCA CCATGGCATGTGTATACCTATGTAACAAACCTGCATGTTCTGCTCATGTATCCCAGAACT TAAAGTATAATAATAAAAAAAAAAGAAAGCACAAAAATAAAAGTACTTGGAAAAGTTTAAA GGGTTAAATATTATGCAAAACTGAAAACTAGCTTCAGATACATTTAAGTTTATATCATGT TAACAAGTTATTTCTTTCTAAAAAATTCTAACCTGTAACACAGAGAGTGGACTTGAACTT GAAAATATGGTTAAGGTACAAATGCAGATTTGGGGTCCCAGTCTCCCAGACTGTGGCTTC TATGGAAGAGATTGTACTGGCTCCAAATTCCACAGATGATTGAACAACTTGTTTCTGCCT GTGTCAGAGCTGAAGAGTGAATATCTCCACTATATATATCTCAAAATCTCCCAAATGAAA TTTGGTAACCCTCTATGCCATAACACATCACATTAATAATTTGTATTCAAAAGTCTCTCA GAAAAGATTTTTGAAATGCCAGATACTTTAATTTTTTTATGTTTATATATTTAGGGTGTA TGAGTACAGATTTCTTACATGCCTATATTGCATAGTGGTGGAGTCTGGGCTTTTACTGTA

FIG. 1AQ

44/51

GTCATCATCTGAACAGTGAACTTGTACCAAATAAGTAATTTTTCAACTCTCATCCACCCA CCCTCCCATCTTTTGTAGTACCCAAGGTCTATTATCCCACTCTGTATGCCTGTGTACCTA TTGTTTAGCTTCCACTTATAAGTGAACACATGCAGCATTTGACTTTCTGTTTCTGAGTTA TTTTACTTAGGATAATGGCCTCCAGTTCCATCTACATGGCTGCAAAAGTTATGATTTTAT TCTTTTTTATGGCTCCATTATATGTATGTGTGTGTATCTCAATTTTCTTTATCAAACCCT CTGTTGATGGACACTTAGATTAGTCCACATTTTTGCTATTGTGATAAACATGTAAGTGCA GGTATCTTTGTAATATAATGATTTCTTTCCCTTTGGATATATACCAGGTAGTGGGATTTC TGGATCTAATGGTAGTTCTATTTTTAGTTCTTTGAGAAATCTCCATACTGTCTTCCATAA AGGTTGTACTAGTTTACATTTCCACCAAAAGTGTATAAGCATTCCCTTTTCTCTGCATCC TCACAAACATCCTTTGCTTATTGACTTTTTAATAACAGCCATTCTGACTAGTGTGAAATA ATATTTTATTGTGATTTTAATTTTCTCTGATGATTAGTGATGTTGAGCATTGTCTCAACA TCACTATGCTAGTGGCATGCATGTTTTCTTTTGAAAAAAAGTTTGTGTTCTTTGCCCACA GAAAATTATTCCTTTGTCAGCTGCATAGTTTACAATTTTTTTCCCATTCTGTAGTTTGTC TGTTCACTCTGTTGATTGTTTATTTTTCTGTCCAGAAACTTTAGTTTAAGTCCCATTTGT CTATTTTTGTTTGTTGCATTTGCCTTTGAGGACTAGGTCATAATTTTTTGCCTGGGCA AATGTCCTGAAGATTTTTTTCCAGGCTTTCTTATAGTATTTTTATAGTTTCGGGTCTTAT TTTCAATCTTCTACATATGGCTATCCAGTTTTCCCAGCACCATTTATTGAATAGGGAGTC ATTTACCCAGTAAATATTTTAGTTGACTTTGTTAAAAATCAGTTGGTTATAGGTGTGTGG TTTTATTTCTAGGTTCTCTATGCTGTTCTATTCATCAATGTGTACATTTTTATACTAGTA CCATGTTGTTTTGGTTACTATAGCTTTGTAGCATAATTTGAAGTCATAATATGATGCCAA CAACTCTGTTCTTTTGTTTGAAATTGCTTTGGCTTTTTTTCCTTGTGAGAGTTTGCTGA GAATGATGGTTTCCAGCTTTGTCCATGTCGCTACAAAGGACATAATCTCACCCTTTTTTA TGGCTGCGTAGTATTCCATGGTGTATATGTGCCACATTTTCTTAATCCAGTCTATCATTG ATGGGGGGGGGGGAAGGGATAGCATTAGGAGATATACCTAATGTAAATGACGAGTTAAT GGGTGCAGCACACATGGCACATGTATACATATGTAGCAAACCTGCACATTGTGCAC TGGCTTTCTGGACTCTTTTTTTTGGTTTTATATGAATTTTAGGATTTTTTTCTAATTCTA TGAAAAATGGCATTGGTAATTTGATAGGGATTGTGTCGAATCAGTAGACTGCTTTAGACA GCATGGTCATTTTAATAATATTGAATCTCTAATCCATGAGCCAGGGATATTTTTCCATTT GTTTTTGTCATCTAGGGTTTTCTTCCATCAGTGTTTTGTAGTTCTCCTTATAGATATCTT TTACCTCTTTGGTGAAATGTATTCCCAGGCATTTTACTTTATCTTATCTTTTTGTAGCTA TTATAAATGGAATTGCTTTCTTAGTTTGGTCCTTGGAAATGCCAACTACATTTAAAATCC TTTTCCATTTGATGGATTTCAGGTCTTGATGAACATCTCAGTTGTAATTTTCTTAAGATT GAAAAAGTAAATATTTTTTCTATATGTATATATAAAATTGTCCTCTCAAAATTTTAAT TCAATAACCTGCTAGATATCACTTTAGAATCTTGCAGTACTAGTTTTCTTCTCAATTAAT TGTAGATCTTAGCCTTTTAATTTGGGCATGTTTTTCCCTATTAGGACTTAAGTTATTAGG ACCTAAGTTTGTAGACAAGAACTATGTTATATTTGAGAAATTTGTGAGTCATGTACTGGG CCTAGCACAGTGCCTCATAAGATGTAGACCCTCAATAAACTTGTTGAATAGGTTAATAAA CACCCAGCCTGGAGTGCAGTGGCACGATATCGGCTCACTGCAAGCTCTGCCTCCTGGGTT CACACCACTCTCCTGCCTCAGCCTCCTGAGTAGCTGGGACTACAGACACCCGCCACCATG CCCGACTAATTTTTGTATTTTAGTAGAGACGGGGTTTCACCTGTGTTAGCCAGGATGG TCTCGATCTCCTCACCTCATGATCTGACCCCCTCGGCCTCCCAAAGTGCTGGGATTACAG GCATGAGCCACCACCTGGCCTATTTCAGTCAATTGTTAAAAGTGCTAAGAACAAGTGG AGATCTTGTTAATGAAGAAAAAAAAAATAGTATTTACTACTTACCTAAACACTCTACTAA AAGAAGAACGTAAACAAAGTCAAAAATGCATTTTTTAGGTGCTAGAGATTAGACAGGACA AAATCTTCTGGCTCTGCCTAGAGTTAAGTGGCTTTGGGAGAGGCTTTGCTGTAGTTTAAA GGCAGAGGTGGGGAAGGCCACTCTGGCCACAAGGACAGATCCACAATGGGATGGGGTATG AAACAGCACGAACCCTTCAGGAAATTACACATAATTTAAAAGGAAAATGGGAGCCCATGG CAGAAAATAGAATTGAACAGCAGGAAAAGGGTAGATAGTAAAAAGCATTTTATAATATTC **AAGGACATTTGAAACTTGTGGTATACAATGAGGAAGAATTTAAAAATTCTATACAGAGGA** GTGACATAGTTAGATTTGTGTTCTGGGGAGCATAATAATAGCATTACAGCGGGTGAATTT GAAAGCTGGGCCTCAAAAGTTTAGATCTCAAATAGGTTTTTATGGGAGTATTCATCCTCA TGAAACATGATTTGGAACTAAACCAAGGCAGTGGCAATGGGGCTGGAAAATAAACACTAG

FIG. 1AR

45/51

ATTTCATATCTAGATGAAGATTTGTGGAATAAGAGAGGCCACATTAATGTTTAATTCTAT TTACAATGGATCCCAGCCACCATCCGCTTTAACACAGAGGTGCTTTTCCAGTAGCTAAGA GGACTAGGTGCTTTAGATACATTTGTGAAGTTGTCCTCCCATTGTTAACATGCTTTTTTT ATTGTCTGTGTGTAGGTTGATGGGGGGGGGGGGGTTAGGATCACACATAGAAGTTCAGTC TTTGAAATGCTTTCTCTTTTTCCCCAAACAATGACCCCCACCTTTTCCTTCTGGCA TATGTTGCCTCAAGACCCTAACACTGCTGCCAATCTGCTGGTCTTAGAGCCAAGAATCTG CCACCACCTGGCCCACCACAGCCTGCTCTGCTAGCTGCTCTCCTGCCAATACTGGCCTTC ATGTACAAGTGTAGGTTTTGAGGGTTCCGTTCTCCCCCTTTCTCTCTTTGAGTGTGGG TTTGTGAGTGTGTGTCTTCTGTAATAAGAAGAAAACAGGCCACATTTTCTCTACTCGT GTTATACACTTCCCGGAGTGTCTCACATCAAAACCTGTCCTAAGTCCAAGCCTTAGAAGC TCTTTGCTGGCCCAGCCTACACTTGGGTTGTTACTTCTCAGGAGCTACCTTTCTGTCAGT TGAGATTTTAACAACCCACCACAGTACTCCAAGCGTGCAGTCCCTCACATCTTGAAATCT GTGCTTTGGCAGCAGCAGAATCAGGGGTCTGTGGATTCTGAACCCAGAATGTGTCAAACC AAAGGGTGACATATTGGGACATTTAATAAGTCAGAGACTATTTCCCAGGAATATTTTTTT GAAGCATTTAAACTAAAATACAATTGAACTGAGATCTCAAAACAGGAAAAATGAACTTGA CAAGAATTAGGCTAAGCTGCATCTCATGACGTAAATATTCACATTTGCATATACATTAAC AGAGTCAAGTCAAAAATTGATTTTTTATTGGATAGGATTAACTTTAGCTACAGAAAACAG AAAGTTTAAATGACTGGCTTTAAAAAAGCAAAAGTTTACTTGCTTTCATTTATATGCAATC TGGGGGGTTGCGGGGAAGATTTGTTTGTGGGTTTCCCATTCTCAAGGTGCCAGGCTCCT GTGTTTCTGCCCCATATCCTTAGAGGGAGGGTTTCCTCCTCAGGATTGCCTTATGTGCAA GGTGACTACTGAAGCTCCATTTTTTATGCCCAAATTGTAGCAAGAAAGGAAAAGGAAGA AGGAACAGAAAGGCACATGACATCACTTCAAATAAAATTAGGGGGAAAATAATAACAGAT TTAAGATCCTTGAAAACAAGGAGTAATAGTTGAGGAAAAGCTTCTTAGCAGCCTGGGACT AAAAACTTCAAAAAATTTAAGATAAAAATCTGAAAACTGGTAGAGAATTGGGGAGAAAAA GAGAATTTGAACAAGGCATGCAAGAGTAAGAAAATGTCATCACAAAATTACTAAGAAAG CATAAAAGCAACTATATTTATTAGAGTAAAAATAAATGGATTGAATAACCCTAGTAAAA ACTAAATTTAAGTGCAAAGATCTCCCAGGGAAGTCAAAGCAAAAATAAAGTCAGATGTTG CTGTCATTAGACAAAGTAAAATTTAAGGTGAAAACATGACAAAGAGGGACATTAAATAAG GATAAATGTACAATGATGGTGACAAACTTTTATAAACTTGAAATTATATAACAAAACA TAAATCATGTAAAACTAAAATACCTTGATAAAATGCAAATTATCAGGTAACAAGAATATA TCTATTGCAGAAAGTAATATATCAAACTAAAATAATGTGTATCTATTACAAGTATACAAT ACTTTGTAGCCTACAAAATAAGAATATACGATTTCTTCTTAAAATGTTATACATTTACAA TAATTAATATTTGGCCACTCAGAAAACCTTGGTAAAGCAAGGAAAGTAGAGATATTATA AGCCAACTTAATAATTTAGTAACATTGGGTAAAAATGGAAGAAGTATCATATTGTGGTTG TGAACATAAGCTCTAGTTCTCCTAGTTTTTGTGATTTGGGAAAGTTAATTATCTTCTCTCT ACCTCGTCTTAATTTTCAGTAATATTAGGATAACAATAGTTTGTACATCATCAGTGTTTT TTTTTTTTGAGGAATAAATGACTCACATGTATTAAACACTTAGATCCATTGTTAACATAT **AATATGTATAAATAATGTCAGTATAAATCAATGTCAGCCTAAAAAGTTAAGACTGTGATT** TTAAATAATACTAGATTTAGAATAAAATCAAAATTGAAATGACATTATTAACTTAAAAAT AACAAAAAAGAGAAGACTTTAAACACAATGGATGGAAAGCAGCTATACCAATAAAAGAC AAACATAGAATTTTAAAAATTTAAATGTTGGAGGGATTAGGTCAGATAAGAGAAATTTCTG TTAGCAGCAGCTGAATTTCTGCTAATAACAGAGAATTGTGAAAAGATGATTTCATAAAT ATGGCAAATGTTTGTAATAGCCATCCTAGGAGCACGGATATTAGTAACTAATTGAGGAAG TACTGTTGGGCAGTGTCAATATACTGGTTAAGAATAGAATTTAAAATAATGCTAATTATA AGGCCAAAAAACTCAGTAATGCAATTTTTTTAGTATAATTCAGTAGGGGAGAAGGAGAGA TAATTAAACTTGGAAATTGACATACAGTTGTCCCTTGGCATCCATGAGGAATTAGTTCCA GGACTCCCTATGGATACCTAAATTCACAAATGCTCAGGTCCCTTATATAAAATGGCAAAA TATTTGCATATAACTTACACACCCCCCTCTTTATAATTTAAGTCATTTCTAAAGTACTTAT AATACCGAATGCATTATAAATGACTGTGGAAATAGTTGTTGTTATTATTTAGGGAATAATG TTGATCCACAGTTGGCTGAATCTATGGATACAGAGCCCACATTTACTGAGGGCAGACTAT ATTTAGAGTACTTAAGGATCACAAGGGACACATCTGAGGGTACTGAAGAGTGGGAAGA AATTACTAACCAGAGGTCAGACTAGAAGGCAAGGAAGTGAAGCCAGGAGATGATTAGAA AATAAGAAAATCATACAAGCCTGGAGATTATGTTGAAGTGTAAGAACATAATTAGAGTGA

FIG. 1AS

GAAACATGAGTCAAGGAAGAAGGAGATTGGTGCTTGAGAGATGTGGCAGACTGTATCTTT CAAAGATGGCTACACCAATATATATCTCATTCCACAAGCTGTTTTTACCATGCTGTATTG TAATTGCCTTGATCAACAGATTGCAGTAGGAGTGATGCTGGATGATTTCAAAGGCTAATA CACACAAGAAAATAATGGCTTTCATTTGACTCTTTCTTGGAACATGTGCCTTGGAAACCA TGAGCTTATTTGCAAGAAGCTCAGCTATCCTAAAGTTTATCTACTGGGTAGACCAAGTGG AGAAATTACACAGACATTGAGATTATGTTCAAGGGGTCTCAGAGGTTCAAGGCCTCCCAA TTCAGGCACCAAACAAGTGGAGAAAAGGCTTTCAAGATCATCCCTCTGAAATAATTGTCT GATTGAAACCTCAAAAGAGTCCCTGAGCCAGAACCATCCAGCCAAGCCACTCTCAAATTC CAAATCCACAGACACCATGAATGACAGTAAATCATTATTGTTGTTTTAAAGCACATAAGT TTTGGGGGGTTATTTACACACAGCAACAGAAAAAAAAACTGATGAATGGGAAACATGGAG AGAAATGCAAATAGAATAAAATGGGAAGGAATACAAGGAGGAGGAAAGTAGTATTGTGCAA AGGTCAGATTAAATTGGAAGGTAACAGGTACAAATATCATTAATGCTAAATTCTATTTGT AATTTATTTAGAAATCTATAGAAGAAGAAAAAGAATAAAAGCCATTGGAAAAGTTTTTAC AATTATTCCATTAAATAGACAAAGTCCTTTAAGGAAAGGGATTAAAATGAAGGTAAGGTG ATCTGCTTAAAAATAATATAGCAATCTGGGAGCCATGGCTCATGCCTGCAATCCCAGTGC TTTGGGAAATCTAGGCAGGAGGACCTCCCAAAGGAGGACTTGGAGTTTGAGACCAGCCTA GAAATAATATTGTATTAATTCCAGTAAAAGCATCAGACCAATTTAGAATATGGATGAGAG AGAAAAACTAGAAATAACACCACAACAAGGAAGGAGAAAGCTGGTCTCTGGCAGGGACTT CTAATTTAGAGAAAGACAGATGATAGCAAACAGCAAAAGTTGTATTATAGATGTAACTTA AAAACTAATTTGATTTTTATTTTTAGTCAGAAAACTGCTTTAGGTATGGAACAAGTATAA CTCTGCTTCTCTTTATAGCAAATTCCCTTGAAAGAGAGACTACCTGGATCAGAAATTCCT CTGCTTCAATTTGATCCTGAATCCACTTCATCTAGATCTTCCTCACCAATTCCCCCAAAT ATTTGTCTTATTATGGTCACATGGGACCTCTACTTTGCTATATCAGTAATTTTGTTCTCA TTTTACTTTTTTGTAGTTAATTACTCCCTTCTCCTTGAAACACTTTCCTTGTTTGGCTTC CAATATCTTCGTGATCTTATATTTTTAATGCGTCCTGCTAGCTTCCCAACTAGGTTTCCT ACTTTCACCTTAATTCCCTATGGTTTATTCTCTACAAGAAAGGAATTATAATCCCTTAAA ${\tt TAAAAAACTAAAGTCCTTACTATATGTCTGTAAATTCCCATAGGATCTGGCCCCACAGCC}$ CCTCTGGCCCCTGTCCATTCTGCCCCAATTCTGCCCGGCCACAGTTGCCCAATAG CTGGTCTGTGAACACATCAAGCACATACTTAATCTCAAGGCTTTTGCAATCATTCTTTTC TCTAGTTGTAATCTCTCATTACTTATTCTGAGTGTCTTGTTTCTGCAGTTGCTTTACTTA CTTGACCTATATAAAATAGTAATTCTTACCCCTACAACTCCATTATGTCCTATCTTCTTT GCCTTGCCTTATGTTTTTTTTTTGGAGTTACAGATACCTGATGTAGATAGTATTTACTTT TTTTATGCTTGCATTAATCACCTAGAATATAAACTCCAAAAGAGGAGCTATTTCTCTTTT ATAATCTATCTAATATATCTTGGATATTTGCTCCCACCTAAATTTCATGTTGAAATGTAA TTCCCTGTGTTGGAGATGGGGTCTGGTGGGAGGTATGTGGATCATGGGGCGGATCCCTCA TGAAGGGCTTGGGCCATCCTTTTGGAGAGAAGTGGGCTCTGGCTCTGACTTCACACGAGA TCTGGTTGTTTAAAAGTGTGCGACAGCTCCCCTGAGCTTCCTCTCACTTGCTCCTGCT TTTGCCATGTGAAGTACCAGCTACTGCTTCATTTTCCACCATGAGTAAAAGATCCCTGAG GCCCTCCCTCAGCAGTACATGTCCCTATGCTTGTTGTGCAGCTGGCAGAACCATGAGCCA ATTAAATCTCTTTTCTTTTAAATTACTCAGTCTCATGTATTTCTTTATAGCAATACAAGG TTGGCTTAATACATATCTCTAAAGCAAAAGCTGGGCCTGGTATGTAATAGGTGTTCAATA AACACTGGGTAGATGTCAAGATGACAGTTTTGTTATTCACATATGGACATGGAAAGGTCT TTGTGGTGCATTGTTAAGGGAGCAAACCAAATTACAGAACACTATATAGAGTAGAGCTGT ATAAAATACATATGGTGTATGTTTATAAATATGTCTAGAAAAATTTGAAAGCTATATATC AAATATCATATCATTTATCTTTAGAAGGCTAATTGCATATTTTCAATTTATTGTTTATAA TTTTTTTTTTTTATCTATTATTATAGGTTACTTGTATAATCACAAAAGACAACTGAATAATTCT TTTTGTCTTCATCAACTTTTATTTTAAGTTCTGGGATACATGTACAGGATGTGCAGGTTT GTTACATAGGTAAACGAGTGGCATGGTGGTTTGCTGCACAGATCGACCCATCACCGAGGT ATTAAGCTCAGCATCCATTAGTTATTCCTCCTGATGCTCTCCCTTCCCCTTGGCCCACCAA

FIG. 1AT

TACACCCTAGTGTATGTTGTTACCCCTCATGTGACCATGTGTTCTCATCATTCAGCTCCC CCATATAAGTAAGAATATGCAGTGTTAGGTTTTCTGTTCCTGTGTTAGTTTGCTGAGGAT ${\tt AACAGGTTCTAGATCCATGTCCCTGCAAAGGACATGCTCTTGTTCCTTTTTATGGG}$ TGCATAGTATTCCTTGGTGTATATGTACCACATGTACAACTAATTTCCACAACAAAAAAT GTACTATTACATGGATATAATGTTTATATTCTCTTCACAGAATTTGAGTCACTTGAATTT CCAATAAGTAAATGTGGAAGGTTGGTGGAAATAGTTAGCTGGAAACTCAGAATTGATATT GCACTATCTATGCATTCAAACAATGATACATATGGTGCATATGTATATATGGCAAAAATC TAAGAAATGTAGCCAAATATTAATATTGCTTACACGTAAGTAGTCAAATCATGGTGGTTT TTTTTTATTTTCTTGATTTTGCAAGAAAATTAATAAAGAGGCTATTTACATTTTAATGTA AACATTTCTGAATTTTTTAAAGATTTCTCAATAGATCTAGGTATTCTTCTTAACCAAATA CTGATACTACCGTTAACCACTTCTGGAAAATTCTGGCAATTGGTCCCTTTGGGGAAGAAC TAGAGGAATCACTACTACACACTTACTGTGGTATTCAGTGCCCTTCCTCAAGGGGAAT TCGCCTATCTTTTTTTCTTAAGTAATATTTTTATCTTTAATAGACAAATAATGGTTGTAT TTATTTACGGGATACAAAGTGACATTTTGATGCAAGCATACCTTGTGGAATGATCAAATC AGGCTAATTAACATATCTGTCATCTCAAATGCTTATCCTTTCTTCATTGTGGGAGCACTT AAAATCAATTCTTTTAGCTATTTGGAAATATAAAATATATTTTTTCTAACTATATTTAC CTCTTTGACCAACATCTCCCCTTTCTTTGTCCATCCTCCTAGCCCAGCCTTTGGTAGCCA TCACTGTACTCTGTATTCTATCACTTTGCCTTTTTAAATTGCACATATAAGAGAGATCA TGCAGTATTTGTTGTCTGTCTGACTTATTTCCTGTAGCAGAATGTCCTTTAGGTTAA TCCATGTTGTCATAAATGACAAAATTTCCTGCCTTTCAAAGGCTGAATAGTATTCCATTG TTTATATATACCACATTGTCAAAATCCATTCATCTGTTGATGGGCATGTAAGTTGTTTTC AAATATTGGCTTTATTAATAATGCGGCAGTGAACGTGGGAGTTCAGACATCTTGTTGACA TACTGATATTAATTCCTTTGACTATATACTCAAAAGTGGAATTGCTGGACTGTGTGGTAA TTTTAGATTTTTAGTAACATTCATACTGTTTTCCAAAATAACTGTATGAATTAACAATAC CATCAACAATGTACAAGGGTTCCCTCTGCTCCACATCCTCATCAACACTTGCTAGTTTTC ATGTTTTCGATAATAGCCAGTCTATCAGGTGTAAGATAATATTTCATTGTGATTTAATTA GCATTTCTTTGATAATCAGAGATTTTGAGCCTTTTTTAATATATCTGTTGACCACTTTTA TGTTTTCCTTTGAGAAATGTGTATTTAAGTCGTCTGCCCATTTTTAATAGGATCATTTGT TTTCTTATTATTGAGGGGTTTGAGTTCCATGCATATTTTAGATACTAGCCTTTTATCCAA TGCGTAATTTGCAAATATTTTCTCCCAATCTGTGGGTTGTCTCTTTAACCTGCTAACTGT TTCCTTTCCTTCCTGCAGAAGCTTTTTAGTTTGATGCAATTCCATTTGTCTATTTTTGCT TCCATTGCCTGTGCTTTTGGGGTTAAGAAATCTCTGCTCGATTACATTTATTGATTTGCG TATATTGAACCAGCCTTGCGTCCCACGGATGAAGCCCACTTGATCATGGTGGATAAGCTT TTTGATGTGCTGCTGGATTCGGTTTGCCAGTATTTTATTGAGGATTTTTTGCATCAATGTT GAATAGTTTCAGAAGGAATGGTACCATTCCTCCTTGTACCTCTGGTAGAATTCGGCTGTG AATCCATCTGGTCCTGGACTCTTTTTGGTTGGTAAACTATTGATTATTGCCACAATTTCA GAGCCTGTTATTGGTCTATTCAGAGATTCAACTTCTTCCTGGTTTAGTCTTGGGAGGGTG TATGTGTCAAGGAATTTATCCATTTCTTCTAGATTTTCTAGTTTATTTGCGTAGAGGTGT TTGTAGTATTCTCTGATGGTAGTTTGTATTTCTGTGGGATTGGTGGTGATATCCCCTTTA TCATTTTTTATTGTGTCTATTTGATTCTTCTCTCTTTTTCTCTTTATTAGTCTTGCTAGC GGTCTATCAATTTTGTTGATCCTTTCAAAAAACCAGCTCCTGAATTCATCCATTTTTTGA AGGGTTTTTTGTGTCTCTATTTCCTTCAGTTCTGCTCTGATTTTAGTTATTTCTTGCCTT CTGCTAGCTTTTGAATGTGTTTGCTCTTGCTTTTCTAGTTCTTTTAATTGTGATGTTAGG GTGTCAGTTTTGGATCTTTCCTGCTTTCTCTTGTGGGCATTTAGTGCTATAAATTTCCCT CTACACACTGCTTTGAATGTGTCCCAGAGATTCTGGTATGTTGTGTCTTTTTTCTCGTTG GTTTCAAAGAACATCTTTATTTCTGCCTTCATTTTGTTATGTACCCAGTAGTCATTCAGG TCTAGTTTGATTGCACCGTGGTCTGAGAGACAGTTTGTTATAATATCTGATCTTATACAT TTGCTGAGGAGAGCTTTACTTCCAACTATGTGGTCAATTTTGGAATAGGTGTGGTGTGGT GCTGAGAAGAATGTATATTCTGTTGATTTCGGGTGGAGAGTTCTGTAGATGTCTATTAGG TCTGCTTGGTGCAGAGCTGAGTTCAATTCCTGGATATCCTTGTTAACTTTCTGTCTCGTT

FIG. 1AU

GATCTGTCTTATGTTGACAGTGGGGTGTTAAAGTCTCCCATTATTATTGTGTGGTAGTCT AAGTCTCTTTGTAGGTCACTCAGGACTTGCTTTATGAATCTGGGTGCTCCTATATTGGGT GCATATATATTAGGATAGTTAGCTCTTCTTGTTCAATTGATCCCTTTACCATTATGTAA TGGCCTTCTTTGTCTCTTTTGATCTTTGTTGGTTTAAAGTCTGTTTTATCAGAGACTAGG ATTGCAACCCCTGCCTTTTTTTGTTTTCCATTTGCTTGGTAGATCTTCCTCCATCCTTTT ACTTTGAGCCTATGTGTCTCTGCACGTGAGATGGGTCTCCTGAATACAGCACACTGAT GGGTCTTGACTCTTTATCCAATTTGCCAGTCTGTGTCTTTTAATTGGAGCATTTAGTCCC TTTACATTTAAAGTTAATATTGTTATGTGTGAATTTGATCCTGTCATTGTAATGTTAGCT GGTTATTTTGTTGTTGGTGGATGCAGTGTCTTCCTAGCCTCTATGGTCTTTACAATTTG AGCTCTTTTAGGGCAGGCCTAGTGGTGACAAAATTTCTCAGCATTTGCTTGTCTGTAAAG GATTTTATTCTCCTTCACTTATGAAGCTTAGTTTGGCTGGATATGAAATTCTGGGTTGA AAATTCTTTTCTTTAAGAATGTTGAATATTGGCCCCCACTCTCTTCTGACTTGTAGAGTT TCTGCCGAGAGATCCGCTGTTAGTCTGATGGGCTTCCCTTTGTGGGTAACCCGACCTTTC TCTCTGGCTGCCCTTAACATTTTTCCTTCATTTCAACTTTGGTGAATCTGACAGTTATG TGTCTTGGAGTTGCTCTTCTCGAGGAGTATCTTTGTGGCATTCTCTGTATTTCCTGAATC TGAATGTTGGCCTTCCTTGCTAGATTGGGGAAGTTCTCCTGGATAATATCCTGGAGAGTG TTTTCCAACTTGCTTCCATTCTCCCCGTCACTTTCAGATACACCAATCAGACGTAGATTT TCTTCCAGTTGATCGCATCGGCTCATGAGGCTTCTGCATTCTTCACGTAGTTCTCGAGCC TTGGCTTTCAGCTCCATCAGCTCCTTTAAGCACTTCTCTGTATTGGTTATTCTAGTTTTA CATTTGTCTAAATTTTTTCAAAGTTTTCCAACTTCTTTGCCTTTGGTTTGAATTTCCTCC TGTAGCTCGGAGTAGTTTTATCGTCTGAAGCCTTCTTCTCTCAACTTGTCAAAGTCATTC TCCATTCAGCTTTGTTCCATTGCTGGTGAGGAGCTGCGTTCCTTTGGAGGAGGAGAGGTG CTCTGCTTTTTAGAGTTTCCAGTTTTTCTGCTCTGTTTTTTCCCCCATCTTTGTGGTTTTA TCTACTTTTGGTCTTTGATGATGGTGATGTACAGATGGGGTTTTGGTGTGGATGTCCTTC CTGTTTGTTAGTTTTCCTTCTAATAGACAGGACCCTCAGCTGCAGGTCTGTTGGAGTTTG CTAGAGGTCCACTCCAGACCCTGTTTGCCTGGGTACCAGCAGCGGTGGCTGCAGAAGAGC GGATTTTCGTGAACCGCGAATGCTGCTGTCTGATCGTTCCTCTGGAAGTTTTGTCTCAGA GGAGTATCCTGCCGTGTGATGTGTCAGTGTGCCCCTACTGGGGGGTGCCTCCCAGTTAGG CTGCTCGGGGGTCAGGGACCCACTTGAGGAGGCAGTTTGCCCGTTCTCAGATC TCCAGCTGCGGGGGGGACACCACTGCTCTCTCAAAGCTGTCGGACAGGGACATTTAA GTCTGCAGAGGTTACTGCTGTCTTTTTTGTTTGTCTGTGCCCTGCCCCCAGAGGTAGAGCC CACAGAGGCAGGCAGGCCTCCTTGAGCTGTGGTGGGCTCCACCCAGTTCGAGCTTCATGG CTGCTTTGTTTACCTAAGCAAGTTTGGGCAATGGCGGGCACCTCTCCCCCAGCCTTGCTG CCACCTTGCAGTTTGATCTCAGACTGCTGTGCTAGCAATCAGCAAGACTCTGTGGGCATA GGCCTCTCCAGCATATAAACAGAACCAAAGACAAAAACCATATGATTATCTCAATAGATG CAGAAAGGGCCTTTGACAAGATTCAACAACGCTTCATGCTAAAAACTCTCAATAAATTAG GTATTGATGGGATGTATCTCAAAATAATAACAGCTACTTATGACAAACCCACAGCCAACA TCATACTGAATAGGCAAAAACTGGAAGCATTCCCTTTGGAAACTGGCACAAGACAGGGAT GCCCTCTCTCACCACTCCTATTCAACATAGTGTTGGAAGTTCTGGCCCAGGCAATTAGGC AGGAGAAGGAAATAAAGGGTATTCGATTAGGAAAAGAGGAAGTCAAATTGTCCCTGTTTG CAGATGACATGGTTGTATATCTAGAAAGCCCCATTATCTCAGTCCAAAATCTCCTTAAGC TGATAAGCAACTTCAGCAAAGTCTCAGGATACAAAATCAATGTACAAAAATCACAAGAAT TATTACACACCAATAACAGACAAATAGAGAGCCAAATCATGAGTGAACTCTCATTCACAA TTGCTTCAAAGAGAATAAAATACCTAGGAATCCAACTTACAAGGGACGTGAAGGACCTCT ACATTCCATGCTCATGGTTAGGAAGAATCAATATCGTGAAAATGGTCATACTGCCCAATG TAATTTATATTCAATGCCATCCCCATCAAGCTACCAATGACTTTCTTCACAGAATTGG AAAAAACTACTTTAAAGTTCATATGGCACCAAAAAAGGGCCCGCATCACCAAGTCAATCC TAAGCCAAAAGAACAAAGCTGGAGGCATCACACTACCTGACTTCAAACTATACTACAAGG CTACAGTAACCAAAACAGCATGGTACTGGTACCAAAACAGAGATATAGCTCAATGGAACA GAACAGAGCCCTCAGAAATAATGCTGCATATCTACAACTATCTGATCTTTGACAAACCTG AGAAAAACAAGCAATGGGGAAAGGATTCCCTATTTAATAAATGGTGCTGGGAAAACTGGT TAGCTATATGTAGAAAGCTGAAACTGGATCCCTTCCTTACAGCTTATTCTAAAATTAACT CAAGATGGATTAAAGACTTAAACGTTAGACCTAAACCATAAAAACCCTAGAAGAAAACCT

FIG. 1AV

49/51

AGGCATTACCATTCAGGACATAGACATGTGCAAGGACTTCATGTCTAAAGCACCAAAAGC AATGGCAACAAAAGCCAAAATTGACAAATGGGATCTAATTAAACTAAAGAGCTTCTGCAC AGCCAAAGAAACTACCATCAGAGTGAGCAGCCAACCTACAAAGTGGGAGAAAATTTTCGC AACCTACTTATCTGACAAAGGGCTAATATCCAGAATCTACAATGAACTAAAGCAAATTTA CAAGAAAAAACAAACCCCCATCAAAAAGTGGGTGAAGGATATAAACAGACACTTCTC AAAAGAAGACATTTGTGCAGCCAAAAAACACATGAAAAAATGCTCATCATCACTGGCCAT CAGAGAAATGCAAATCAAAACCACAATAAGATACCATCTCACACCACTTAGAATGGCAAT CATTAAAAAGTCAGGAAACAACAGGTGCTGGAGAAGATGTGGAGAAATAGAAACACTTTT ACACTGTTGGTGGGACTGTAAACTAGTTCAACCATTGTGGAAGTCAGTGTGGCGATTCCT CAGGGATCTAGAACTAGAAATACCATTTGACCCAGCCATCCCATTACTGGGTATATACCC AAAGGACTATAAATCATGCTGCTATAAAGACACATGCACACGTATGTTTATTGTGGCACT ATTCACAATAGCAAAGACTTGGAACCAACCCAAATGTCCAACAATGATAGACTGGATTAA GAAAATGTGGCACATATACACCATGGAATACTATGCAGCCATAAAAAAGGATGAGTTCAT GTCCTTTGTAGGGACATGGATGAAATTGGAAATCATCATTCTCAGTAAACTATTGTAAGA ACAAAAACCAAACACCGCATATGCTCACTCATAGGTGGGAATTGAACAATGAGAACACA ATAGCCTTAGGAAATATACCTAATTATAAATGACGAGTTAATGGGTGCAGCACAGCAGCA TGGCACATGTATGCATATGTAACTAACCTGCACATTGTGCACATGTACCCTAAAACTTAA AAAAATAAAAAGAAATCTCTGCTCATATCCAGGCCATGATGGTTTCCCCCTGTGTTTTCT TCAAGTAGTTTTATAGCTTCAAGTCTTATGTTATATTAAGTCTTTAATCCATTTTGAGGT GTTTTCCCAACACCATTTATTGAGAAGTCTGTCATTTCCCCATGGTGTGATCTTGTTACC TTTATGAAAATTTAATTGACCATAGGTGTATGGGTTTATTTCTGGGCTTTCTATCATATT ${\tt CCATTGATTGATATGTCTGGTTTTATGCCAGTACTATGCTGCTTTGATTACTGTGGATTT}$ GTAATGTAATTTAATGTCTGAGAGTGTGAAGCCTGCAGCATTATTTTTTCTCAAGATTGT ${\tt TATCTGTGGCTATTTGTAGTCTTTTGTGGTTTCATATATTTTACAATTTTTATTTCT}$ GTGAAAAATGCATTGGAATTTTCATATGGATTACATTTAATCCGCTTTGGGTAGTATGAC GTCATCTTCAGGTTTTTTCAACAATGTTTTATAGTTTTAGTATATGGATCTTTCACTTCC GTGTGTGTGCATCAACTAACCATAGTCATGTGGGTTTATTTCTGGGCTTTCTATCATGTT ACTGTTGTAATTTTAAAATTTCTTTCTCAGGTTGTATGTTGTTAGTGTACAGAAATAATA TTAATTTTGTAAGTTGATTTTGTATTCTGCAAATTCACTAAATTTGTTAATTTGTTTTAA AATTTCATTTATTCTTTTCCTATTTGGATGCTTTTTATTCTTACCCAATTGTTTTGACTA ${\tt GGACCTCCAGTACTATGTTGAACATAATTGATGAAAGCAGACATCCTTGTCTTGCTCCTG}$ ATCCAAAAGCCTTTAACTTTTCACCACTGAGTATGATGTTCACTGTAGGCTTGTTATATA TGGTCTTTGTTGTGCTGAGAAACATTCCTTCTATAACTGATTTTCAAAAGTTTATCATGA AAGGATGTTAAATTATTTCAAATGTTTTTTTCTTCATCTATTGAGGTGATTATATTGTTTT TATTCTTCATTCTGTTACTATGGTGAATCATATTTTTAATTGTTTTTTACTTGCATAAAT TTATTTTGTGATAGGTAGAAAAGCACATCTGCAGACCTAGAAGCAGAGTGAATCTAAAAA ATATTATTATAATTATTATGAGTACACAATAGGTATATATTTTCATGGGGTACATTCAA TGTTCTGATACAGGCATATGATGTGTAATAATCACATCAGGGTATTTGGAGTATTCATTA CCTCAAGCATTTATCATTTCTTTGTGTTAGGGAATTTCAGTTTCATTCTTCTAGTTATTT AAAATATACAATGAATTATTATTGACTGTAGTCACCCTGTTGTGCTATCAAATAGTATGT CTTATTCATTTTATTTAACTATATTTTTGCACCCATTAACAATCCCCACTTGATTTGAAT ATGGTAAGCCATTCTTGCATCCTAGGAATAAATTCCATTTGACCATGGTGAATGATCCTT TTAATGTACTGTTGAATATAGTTTTTGGTATTTTGTTGAGGATTTTTGCATCCATGTTCA TTGGTGTGCTGCACCCATTAACTCATCATTTAACATTATGGAAAATCTCCTAATGCTATC CCTCCCCGCTCCCCCACCCCACACAGGCCCCGGTGTGTGATGTTCGCCTTCCTGTGTC CATGTGTTCTCATTGTTCAATTCCCACCTATGAGTGAAAACACACGGTGTTTCTTAGTCT GGCTTTGGTCTCAGGCTAATGTTGGCCTTACAAAATGATTGTGGAAATATTTCCTTCTCT TCAATTTTTTGAAGAAGTTTGAAAATAATTATTACCAGTTCTTCTATAAATGTTGGGTAG

FIG. 1AW

AATTCATTTATGAAAATATCTTTTCCTGGGTTTTCCTTGATGGCGGACTTTTCATTACTG ATTTAATTTCCTTGCTCATTACTGTTCCATTTATATTCCTCATGATTTGATCTTGGAAGG TTCGTAGTGGTCTCATAAGATCCTTTGTATTTTTGTACTATCAATTGTGATATCTTTTTT CATTTCTGCTTTAGTTTACTTGAACCACCTGTATTTTCTCGTGGTTAATTTAGCTAAGGA TTGTCAATTTTGTTTTTTTGGAAGACCAACGCTTAGCTTTACTGATCTCTTGTATT GTTTTTCTAATTTCTATTTCATTGATTTTTGCTCTGAAATGTTTCCTTTCTTCCACTAAC TTTAGGCTTAGATTGTTCTTTTTACTAATTCATTGAGGAGTAACATTAAGTTGTTTAT ${\tt TTAAGATCTCTCTCTCTCTCTCTCTCTCTCTCTTTTGATGTAGGCATTTAGTGTTACAAA}$ CTTTCCTCTTAGAACTGCTTTTGCTGAATCCTGTAAGTTTTAATATGTTGTTTCCATTTT CATTTTTCTCTAAATATTTTTAAAATTAATTTTTGAATTTCCTCTTTGACTCAATAGTTT TTCAGGAGCATGTTTAATTTGCATATACTTGTTAATTTTCTTGGTTTCTCCTGTTA TTGATCTATAGCTTTATATCATTGTGATTGAGAAAGATACTTGATATAATGTTGATCTTC TGACACTTGTTAAGATGTTTTGTGGTCTATCAATTGATTTATCCTAGTGAATGTTACATG TATACTTGAGAAAAATGTATATTTTGTTGCTGTTGGATGAAATGTTCTGTATAGGTCTAT TAACTCCATTGGTATACGTATAGTTCAAGTCATATTTTGTTATTAAAAATTTTTTGTCTA GATAATAGTTCTGTTGGTTGGAAGTGGGATATTAAAATTATTTACTATTATTGTGCTGCAT TTATGTCTCTTTTCAGAACTCTTAATCTTTGATTTATATATTTAGGTGCTTCAGTGTTGG GTGCATATATTTACAATTGTTATATTATCTTGATGCACTGATCTTTTTATTATAATAT ACTGACCTTCTTTATCTCTTTTTACAGTTTTTTTTAACCTAAAGTTTATTTGGTGTGAAA TAAGTATAGCCACCCCTGCTCTGTTTTATTTGCCTGGAATATCATTTTCCATCACTTCAT $\tt TTTCAACCTGTAAGTTTCCTTTAAGGTAAGGTGAGTCTTCTGTAGGCCCATATAGTTGGA$ TCTTGTTTGGTATGTATCATGGTACTGTATGCCTTTTGACTACAGAATCTAATCCATTAA ACTTTAAAGTAATTATTGATAGATGAGAGGTTGCTACTTCCATTTTATTGTTTTCAAGTT CTTTTTGCAGGGATATATTTTGAATTTTTTAAAATATTTTGTGTATCTATTATAGGCTCA TGCTTTGTGGTTACATAAATCATCTTATACCTATAACAAGCTATGCCAAGTTGATAACAA CTTAAGTTTGATCACTTACACAAAGGCTACACTTTTACTCTCCTCCTTCTAAATTTTATG TTTTTGATGTCATTCTTTACATCTTTTTATAATATGCATACTTAACAAACTACTGTAGCT GTAGTTGCTTTTAAGAATTTTGCCTTTTAACCCTTATACTAGAGAAATCCTTGATTTGTT CACCATCATTACAATATTAGAATGTTTTGGAATTGAAAAATGCCATTAATTTTACCAGTG CGTTTTATACTTTCATATGTTTTCATGTTTCTATTTTGAATCCTTTTCCTTCAGCTTGAA GAACTCCCTTTAGCATTTCTTATAACGCAGGTCTAATGGTGAGAAACTCAGCCTTTGTTA CTCTGAGAAAGTCTTTAACATCCCTCATTATTTAAAGACAGGTTTGCTAGGTATACTATT CTTGATTGGCAGGTTTTTTTTTTTTAGAATTTTGAATATATTATCCCACTCCCTTGAGCT TTCAAGGTTAATGCTGAGAAATTTGCTGATAGTTTTATCAGGGTTCTCTTATATGTGACA ATTCAATTCTCCCTTGCTGCTTTCCATACTCTAAGTTTTGACAGTTTTGTTATGATGTGC CTTGGTGTGAGTTTCTTTTCCTTTTTTAAATTTTAGATTCAGAGGGTACATGTGCAGATT TGCTGCAAGGACATATTGTGTGGCGTTGGGCTTCTGTTGATCCCACCACTCAAGTGGTGA CTTTTTCCCTTTTTGGAAGATGCAGTGTCAATTGTTTCTATATTTATGTCTGTGTGTACC CAATATTTAGTTCCTACTTATGTGAAAGAACATGCAATATTTGGTTTTCTGTTTCCGTGT TAATTTGCATAGGATAATATTTTCCAGTAGTCTGTCCATGTTGCTGAAAAAGACATGAGT TTGTTCTTTTTTATGGCTTCACAGTATTTCATGATGTATATGTACTTGGTGTGGATTTAT CCGGATTCATTTATTTGGTATTCTTTGGGATTCCTGTATCTGGCTTTCTATTTTCTTCC CCAGTACTGGGAAATTTTCTGCCATTATTTTTTGAATATGTTCTGTGCTTGTCTCTCT CCTCCTTCTGAACACCTATAATGTATATTTGCTCTGATTGAGGGTGTCAGTATGTCTCT TAAGATGTGTTCATTCTTTTCATTCTTTTTCCTTTTTGCTGCTTAGATTGGATGATTTC CAGTGACTTGTCTTTGAGTTCATTGATATTTTCTTCTGCTTAATCTCATTTGTGGGTGAA CTTTCATATACTTTCTCTTTGTTAAAGTTCTCTGTTTTTTGCATTTCTCTCTGGACCTTAG TGACAGTCTTTATAATCATTATTTTAAATTCTCTATTGGGTAAATTACATCTCTTCTATT TCTTTAGTTTCCTTGACTCTGTGTTGTTTACTGCACATTAGATAAGACAGCTGCCTTTCC CAGTCTTATCAAACAGGACCTGTGTAGAAGAAAAATATCACTAGTCCATTTGACAAAAAA TTTTAATGTGCCTCTCAAAGCTTTGTTTGTCCAGGCCACTGTTTCTGTTATTGGTGGCTC CCAGGAGATTGGGATATGCCATGTCCTATCAATACTCTGTGAACTATAAGATAGAGGCCA

FIG. 1AX

GACTTTCAAAATGTAGCCAGAAAAATGTCAAGTATTAGATGTGTGGTCCAGTTCCTTCTA AGAGGTACTATGGAAACTGCCTGTATTTGTGTTCAGGCCACACTTTTTGATTCTGGGAAG ATAGCTTTGGGAGTGGGGCCACTGTTTGTCTACATCTTTGTTATCTGTGATCTAGAGTAA GTTAGGAATGCAAAGCTCCACCACTCCCAAGCTTAGGCTGTTAAGAATTCAGTCCTTTGG GTGGGAGCTGTAGAAGTTGTGACACTTAATTGTGAACAAACTCTTTTCAAGAAGAATAGG TAGGCTATAAAATAATAGAAGAAATGAATAGAGCTATAGAAGTTGTGACACTTGGTATGT GAACAAACTCCTTTTAGGAAAAATAGGCTGGGGACAAGCCAAGTTCTGCTTAGTCTACCT GAGAGCTACTATTAGTCTGTCTTGTTAGCTCCCTGATGCAAGCTGGAGGTTAAGCTATGT ATTCTAGCCCCTGTTCTCCACTGCTCCCAAGAGATATAGTTCCTGGAAGAGTTTGCATGC CTGTTTAAAACCACCTCTTTGTTCTGTGATCTAGGGAGACTTGTATATGCCTAGTCTCTT CTGCTCTTAGAGCCAGGAGTTTTGGGATATAGTATTTCTGGTAAATGCTGTAAAAGGGCA TTTTGTGGGTGAACACACTCCTTCCAGGGAGAATTGGGAGAGCTGGGATTATTGCTGAGT TGAGCTGGAGGAAGTCTCAGGAAGTGTTAAGCTGCTCAGGCTGTTAGAGAGCTACTT TTTGCTTGCCCCTTTAACTCTCAGATGCGTTAGTTAGAAACCAGACTGTCAAGTAGCCGC TAGGGGAGTATGCTGTAAACCTCTTCCAGGGAGAACCAGGTAGTGGTATTTTTGAGTCCT GTCTCTGTACTAATTCTACTAATTCACAGTGTTAAAGCACCTGAAAAAGTGCTTGCACAC TCTAACTCCCAGAGTTGGTGAATTAAGAGCCAAACTGTTGGGCATCTTATAATTGGGGTG CCATATGTAAGGTCCCAATCCTCTCCACAGGGAGAATCTGAGTGTTAGTGATTCCAGTTA TATGGTGAAGTACCTGGAAGGGGTCCATGCTCAAGTATGCCTCAGATTTGTCTACCCATT GAGACAGAGTCTCATTCTGTTGTAAAGGCTAGAGTGCAGTGGCACAATCTTGGCTCACTG CAGCCTCTGCCTCCCTGGTTCAAGTGATTCTCCTGCCTCAGCCTCCCGAGTAGCTGTGAC TACAGATGCGTACCACCATTCCCAGCTAATTTTTGTATTTTTGGTAGAGACAGGGTTTCA TCATGTTATCCAGGCTGGTCTCAAACTCCTGGACTCAAATAATCCACCAGCCTTGGCCTC GATGTTCATTCTGTGCATTTGTGAGTATTGGGAGTGCCAGGAGCTTCCTATTCTGCCATG TTGCTGACATCAGTCTAAGGAAAACAGTTTAAAGAAAGTTCATCAAAAAGTAACAGTAGA CTTTCTTTCTTTCCTTTCTTTCTTTCCTTTCC

FIG. 1AY